

NON-NUCLEOTIDE COMPOSITIONS AND METHOD FOR TREATING PAIN

TECHNICAL FIELD

This invention relates to a method of treating pain. More particularly, the present
5 invention relates to a method of treating pain with non-nucleotide P2X receptor antagonists.

BACKGROUND OF THE INVENTION

The general term "pain" is defined here to represent all categories including: traumatic
pain resulting from tissue injury, post-surgical pain, burn pain, inflammatory pain; pain
10 associated with disease such as cancer, AIDS, arthritis, herpes, migraine; pain associated with
nerve damage or neuropathy, such as diabetic neuropathy, causalgia, brachial plexus avulsion,
occipital neuralgia, fibromyalgia, gout, and other forms of neuralgic, neuropathic and
idiopathic pain syndromes; and specific organ or tissue pain, such as ocular and corneal pain,
bone pain, heart pain, skin pain, visceral (kidney, gall bladder, gastrointestinal, etc.) pain,
15 joint pain, dental pain and muscle pain. The term "pain" also includes pain of varying
severity, i.e. mild, moderate and severe pain, as well as acute and chronic pain.

In traumatic or nociceptive pain, an external stimulus causes a normal sensory
response to tissue damage associated with an insult or illness, and this sensation of pain is
generally responsive to narcotic analgesics such as morphine; whereas in neuropathic pain,
20 which results from injury to the nervous system, the sensation of pain is typically not
responsive to narcotic analgesics. Neuropathic pain often involves neural hypersensitivity
and can persist without any overt external stimulus (Goodman & Gilman's "The
Pharmacologic Basis of Therapeutics", 1996, 9th Ed., p. 529, McGraw-Hill).

The therapeutic objective of most pain therapy is to alleviate the symptoms of pain
25 regardless of the cause. Current therapies include the use of opioid narcotic analgesics such
as morphine and fentanyl, nonsteroidal anti-inflammatory drugs (NSAIDS) such as aspirin,
ibuprofen and cyclooxygenase inhibitors, or ion channel blockers such as lidocaine and
novacaine. However, these therapies all have limitations. Opioids, for example, can cause
tolerance, dependence, constipation, respiratory depression and sedation. NSAIDS can have
30 gastrointestinal side effects and can increase bleeding time; in general, they are not effective
in treating severe pain. Central nervous system (CNS) and cardiovascular side effects
(Goodman & Gilman's "The Pharmacologic Basis of Therapeutics", 9th Ed., 1996, p. 337,
McGraw-Hill), as well as corneal damage are reported for the non-selective sodium channel

blockers. Given these drawbacks to current therapies, there is clearly a need for better pain treatments.

Numerous published studies have documented the involvement of P2X nucleotide receptors in nociception. Activation of P2X receptors with agonists such as ATP or benzoylbenzoyl-ATP is associated with hyperalgesic action (Chizh and Illes, *Pharmacol. Rev.* 53:553-568 (2000); Jarvis and Kowaluk, *Drug Development Res.* 52:220-231 (2001)). Antagonism of P2X₃ receptors by the ATP analog, ATP-TNP, has been shown to produce antinociceptive effects in rats (Jarvis and Kowaluk, 2001).

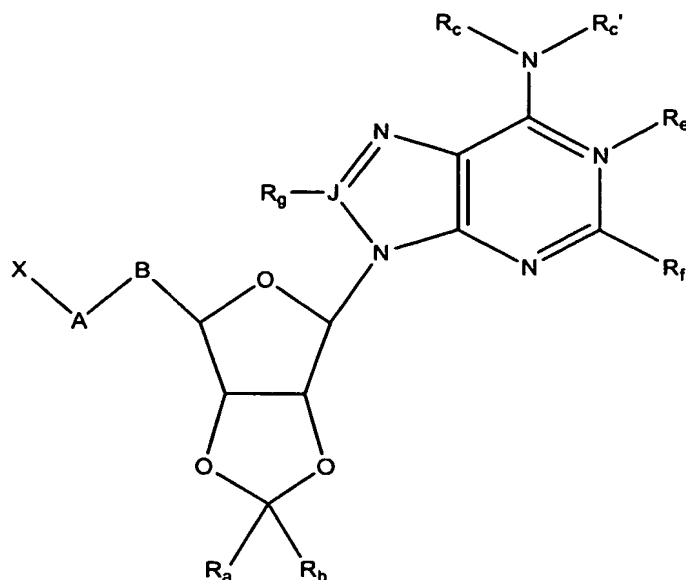
As described above, agents commonly used to treat pain may cause adverse side effects, and thus there is a continuing need for new agents that are both safe and effective in treating pain.

SUMMARY OF THE INVENTION

The present invention is directed to a method of treating pain. The method comprises administering to a subject a pharmaceutical composition comprising an effective amount of one or more non-nucleotide P2X receptor antagonist. The methods of the present invention are useful in reducing pain such as traumatic pain, neuropathic pain, organ pain and pain associated with diseases.

The non-nucleotide P2X receptor antagonists useful for this invention comprise compounds of general Formula I, and/or tautomers thereof, and/or pharmaceutically-acceptable hydrates, solvates, and/or salts thereof.

Formula I



5 The invention also provides novel pharmaceutical compositions comprising compounds of Formula I. The compounds of Formula I are useful in that they possess antagonist activity at P2X receptors.

The compounds of the present method can be used alone to treat pain, or the compounds of the present method can also be used in conjunction with other therapeutic
 10 agents or adjunctive therapies useful for treating pain, thus providing a beneficial therapeutic effect at lower concentrations of each pain reliever than when each is used separately.

DETAILED DESCRIPTION OF THE INVENTION

The inventors have unexpectedly discovered that by administering a non-nucleotide
 15 P2X receptor antagonist to a subject in need of pain treatment, the severity and/or incidence of pain sensation is reduced.

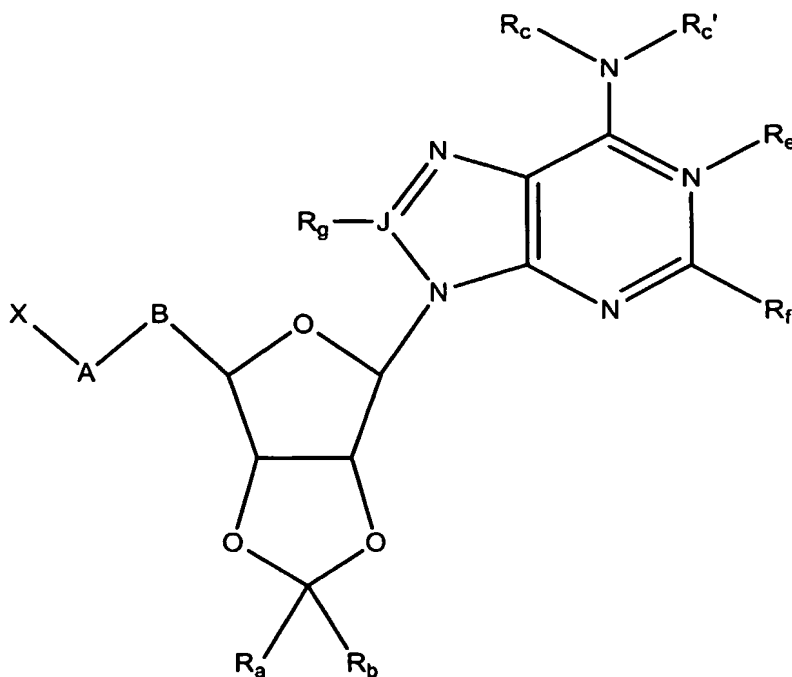
The present invention is directed to a method of treating pain in a subject in need of such treatment, regardless of the cause or location of the pain. The method comprises administering to a subject a pharmaceutical composition comprising an effective amount of a
 20 P2X receptor antagonist, preferably an antagonist of a P2X₃, P2X_{2/3}, P2X₄, or P2X₇ receptor, and more preferably as an antagonist of a P2X₃ or P2X_{2/3} receptor. The methods of the present invention are useful in the treatment of traumatic pain particularly when the pain is associated with tissue damage, neuropathic pain, organ pain or pain associated with diseases.

An effective amount of a P2X receptor antagonist is an amount that antagonizes the interaction of a P2X receptor with another agonist such as ATP, leading to a reduction of nociception and ameliorating the symptoms of pain. The effect of treating pain can be measured in a method such as, but not limited to, abdominal constriction (Collier, *et al.*, *Br. J. Pharmacol. Chemother.* 32: 295-310 (1968)).

The present invention provides a method to alleviate the symptoms of pain. The general term "pain" treatable by the present method includes traumatic pain, neuropathic pain, organ pain, and pain associated with diseases. Traumatic pain includes pain resulting from injury, post-surgical pain, burn pain and inflammatory pain. Neuropathic pain includes, but is not limited to, neuropathic and idiopathic pain syndromes, and pain associated with neuropathy such as diabetic neuropathy, causalgia, brachial plexus avulsion, occipital neuralgia, fibromyalgia, gout, and other forms of neuralgia. Organ pain includes, but is not limited to, ocular, corneal, bone, heart, skin, visceral (kidney, gall bladder, gastrointestinal, etc.), joint, dental and muscle pain. Pain associated with diseases includes, but is not limited to, pain associated with cancer, AIDS, arthritis, herpes, sickle cell anemia and migraine. The present invention reduces pain of varying severity, (i.e. mild, moderate and severe pain) as well as both acute and chronic pain.

The P2X receptor antagonists useful for this invention include compounds of Formula I, or a tautomer thereof, or a pharmaceutically-acceptable salt, -hydrate, or -solvate thereof:

Formula I



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wherein R_a and R_b are each independently selected from the group consisting of: hydrogen, (saturated or unsaturated) C_{1-8} alkyl, (saturated or unsaturated) C_{3-7} cycloalkyl, aralkyl (including saturation and/or unsaturation in the alkylene portion), aryl, and (saturated or unsaturated) C_{2-6} heterocycle; or

10 R_a and R_b can be taken together to form a ring of 3 to 7 members, with or without substitution, and with or without heteroatoms in place of ring carbon atoms;

R_c and R_c' are independently selected from the group consisting of: H, OR, (saturated or unsaturated) C_{1-8} alkyl, (saturated or unsaturated) C_{3-7} cycloalkyl, aralkyl (including saturation and/or unsaturation in the alkylene portion), aryl, (saturated or unsaturated) heterocycle, and -

15 $C(G)\Sigma$; wherein $G = O, S$ or NR_d ; and

$\Sigma = L, R_d, OR_d$, or $N(R_d)_2$; except that $-NR_cR_c'$ cannot be $-N(OR)_2$; and OR_d cannot be $-OH$; each R_d is independently selected from the group consisting of: H, (saturated or unsaturated) C_{1-8} alkyl, (saturated or unsaturated) C_{3-7} cycloalkyl, aralkyl (including saturation and/or

unsaturation in the alkylene portion), aryl, heteroaryl, and (saturated or unsaturated) C₂₋₆ heterocycle; or

two R_d groups can be taken together to form a ring of 4 to 7 members, with or without unsaturation and with or without heteroatoms in place of ring-carbon units; or

- 5 one R_d and one of R_c or R_c' can be taken together to form a ring of 4 to 7 members, with or without unsaturation and with or without heteroatoms in place of ring-carbon units;

R is selected from the group consisting of: H, (saturated or unsaturated) C₁₋₈ alkyl, (saturated or unsaturated) C₃₋₇ cycloalkyl, aryl, aralkyl (including saturation and/or unsaturation in the alkylene portion), heteroaryl, and (saturated or unsaturated) C₂₋₆ heterocycle;

- 10 L is selected from the group consisting of: H, -CF₃, -CF₂CF₃, (saturated or unsaturated) C₁₋₈ alkyl, (saturated or unsaturated) C₃₋₇ cycloalkyl, aryl, aralkyl (including saturation and/or unsaturation in the alkylene portion), heteroaryl, (saturated or unsaturated) C₂₋₆ heterocycle, (saturated or unsaturated) C₁₋₆ alkoxy, aralkoxy, aryloxy, N,N-disubstituted-amino, N-substituted amino, and unsubstituted-amino;

- 15 when L is N-substituted-amino, or N,N-disubstituted-amino, each substituent of said amino group of L is selected from the group consisting of: C₁₋₈ alkyl, C₃₋₇ cycloalkyl, aryl, aralkyl, heteroaryl, and (saturated or unsaturated) C₂₋₆ heterocycle;

when L is N,N-disubstituted-amino, the two substituents independently selected from the group above can then be taken together to form a ring of 3 to 7 members, wherein said

- 20 formed ring thereon bears the remaining features of said selected substituents before said ring formation;

R_c = O or absent;

R_f = H, halogen, (saturated or unsaturated) C₁₋₈ alkyl, (saturated or unsaturated) C₃₋₇ cycloalkyl, aryl, aralkyl (including saturation and/or unsaturation in the alkylene portion),

- 25 heteroaryl, (saturated or unsaturated) C₂₋₆ heterocycle, -OH, (saturated or unsaturated) C₁₋₆ alkoxy, aryloxy, -SH, C₁₋₆ thioalkyl, thioaryl, -[(CO)OR], -[(CO)NRR], amino, -N-substituted amino, or N,N-disubstituted amino; wherein each said substituent on said N-substituted-amino-group, or N,N-disubstituted-amino-group of R_f is independently selected from the group consisting of: C₁₋₈ alkyl, C₃₋₇ cycloalkyl, aryl, aralkyl, heteroaryl, C₂₋₆ heterocycle, -
30 [(CO)R] and -[(CO)-NRR]; wherein each R is independently as defined above; or when R_f is -[(CO)NRR], -[NH(CO)NRR], -[N(C₁₋₈ alkyl)(CO)NRR], -[N(aryl)(CO)NRR], or [N(aralkyl)(CO)NRR], the R groups of a said -NRR unit (N,N-disubstituted-amino group) in

R_f can be taken together such that a ring of 3 to 7 members is formed, with or without heteroatoms in place of the ring-carbon units;

J = N or C, with the proviso that when J = N, then R_g is absent;

when J = C, R_g is selected from the group consisting of: -H, halogen, (saturated or

5 unsaturated) C₁₋₈ alkyl, (saturated or unsaturated) C₃₋₇ cycloalkyl, aralkyl (including saturation and/or unsaturation in the alkylene portion), aryl, -OH, (saturated or unsaturated) C₁₋₆ alkoxy, aryloxy, -SH, C₁₋₆ thioalkyl, thioaryl, -[(CO)OR], -[(CO)NRR], and -NRR; wherein each R is independently as defined above; or

10 when R_g is -[(CO)NRR] or -NRR, the R groups of said -NRR unit (N,N-disubstituted-amino group) in R_g can be taken together such that a ring of 3 to 7 members is formed, with or without heteroatoms in place of the ring-carbon units;

A and B are each independently selected from the group consisting of: -C₁₋₃ alkylene-, -CF₂-, and -(CO)-; wherein each said -C₁₋₃ alkylene- unit of A and B independently can be saturated or unsaturated, and each carbon of a -C₁₋₃ alkylene- unit of B independently can be substituted
15 with 0 to 2 fluorine groups, 0 to 1 methyl groups, 0 to 2 -[(CO)OR] groups, and 0 to 1 -(OR) groups; or

B is absent; or

one substitution of a heteroatom-containing-unit selected from the group: -O-,

20 -S-, -NR-, -[NR(CO)]- or -N[(CO)L]-, where each R and L is independently as defined above, can be made for any one-carbon-unit within either or both of said C₁₋₃ alkylene units of A and B, provided that fewer than three said heteroatom-containing-unit for one-carbon-unit substitutions on the -A-B- chain are made, no -S-S- or -O-O- bonds are formed in the X-A-B- chain by said substitution or substitutions of a heteroatom-containing-unit for a one-carbon-unit on the -A-B- chain, and no said heteroatom substitution is made such that the said
25 replacement heteroatom connects directly to the tetrahydrofuran ring shown in Formula I;

X = -OR, -SR, -S(O)L, -S(O₂)L, -SO₃H, -S(O₂)NRR, -S(O₂)NR(CO)L, -NRR, -NR(CO)L, -N[(CO)L]₂, -NR(SO₂)L, -NR(CO)NR(SO₂)L, -NR(SO₂)NRR, or -NR(SO₂)NR(CO)L;

wherein each R and L is independently as defined above;

30 wherein the R groups of a -NRR unit (N,N-disubstituted-amino group) in X can be taken together such that a ring of 3 to 7 members is formed, with or without heteroatoms in place of the ring-carbon units;

with the proviso that no compound in Formula I contains: a halogen-group, hydroxy-group, sulfhydryl-group, or amino-group (-NH₂, N-substituted-amino, or N,N-disubstituted-amino)

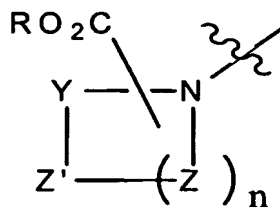
attached to an sp^3 -hybridized-carbon atom that is bonded directly to a heteroatom selected from the group consisting of O, S and N, as compounds in this class (e.g., $-\text{C}(\text{OH})(\text{SR})-$, $-\text{CCl}(\text{NRR})-$, etc.), in general have lower chemical stability;

the first exception to this proviso is: compounds in which the said sp^3 -hybridized-carbon atom is bonded directly to: 1) a sulfur atom which is part of a $-\text{S}(\text{O})-$ -group (sulfinyl group), or a $-\text{S}(\text{O}_2)-$ -group (sulfonyl group), and also to: 2) one or more halogen groups; (an example of a moiety having this arrangement is the trifluoromethanesulfonyl group);

the second and final exception to this proviso is the C-1' position of the furanose of compounds of Formula I wherein the sp^3 -hybridized carbon atom at the 1'-position is attached to: 1) the oxygen atom of the furanose ring and to: 2) the nitrogen atom of the adenine or 8-azaadenine moiety; or

X is a group as provided in Formula II:

Formula II



wherein:

$n = 1$ to 4 , inclusive;

Y, Z and Z' are independently selected from $-\text{CRR}_f$, $-\text{NR}-$, $-\text{N}(\text{CO})\text{L}-$, $-\text{O}-$ and $-\text{S}-$; or the said $-\text{Y}-\text{Z}'$ -unit, taken together, can be selected to be a $-\text{N}=\text{N}-$ unit or a $-\text{CR}=\text{CR}_f$ unit; or any $-(\text{Z})_2$ -unit or subunit of $-(\text{Z})_n$ can be selected to be a $-\text{CR}=\text{CR}_f$ unit;

and

with the provisos that the ring shown in Formula II contains no more than three heteroatoms,

and that the shown pendant $-\text{CO}_2\text{R}$ unit in Formula II is a substituent on the ring described in Formula II, and that the ring of Formula II contains no halogen-group, hydroxy-group, sulfhydryl-group, or amino-group ($-\text{NH}_2$, N-substituted-amino, or N, N-disubstituted-amino) attached to an sp^3 -hybridized-carbon atom that is bonded directly to a heteroatom selected

from the group consisting of O, S, and N, as such types of compounds are unstable in many cases.

Preferably, the furanosyl moiety in Formula I has the 2'- and 3'-oxygen-groups in a cis-orientation, relative to one another on the furanose ring. Further, a furanosyl moiety which supports a 2', 3'-acetal or -ketal group is, preferably, derived from ribose; other furanose derivatives can be used, however. A preferred stereochemical embodiment of this invention includes, but is not limited to (D)-ribose-(2', 3'-acetal or -ketal) compounds of Formula I, such as found in acetals derived from (-)-adenosine.

In one embodiment of the method, the compound of Formula I is selected from the group consisting of: 6-[6-(3-Cyclopropyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxole-4-carboxylic acid; 6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxole-4-carboxylic acid; 6-[6-(3-Butyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxole-4-carboxylic acid; 6-[6-(3-Propyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxole-4-carboxylic acid; 2-[(6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethyl)-amino)-methyl]-cyclopropanecarboxylic acid; 3-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethyl}-3H-[1,2,3]triazole-4-carboxylic acid; 3-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-propionic acid; {6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-acetic acid; 1-{6-[6-(3-Cyclopentyl-ureido)-purin-9-yl]-2-phenyl-tetrahydro-furo[3,4-d][1,3]dioxole-4-carbonyl}-pyrrolidine-2-carboxylic acid; {6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-phenyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-acetic acid; {6-[6-(3-Cyclopentyl-ureido)-purin-9-yl]-2-phenyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-acetic acid; 3-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-phenyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-propionic acid; 3-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxymethyl}-benzoic acid; ({6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethyl)-amino)-acetic acid; 1-Ethyl-3-[9-(6-hydroxymethyl-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl)-8-iodo-9H-purin-6-yl]-urea; 1-Ethyl-3-{9-[6-(2-hydroxy-ethoxymethyl)-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl]-8-iodo-9H-purin-6-yl}-urea; 1-Ethyl-3-{9-[6-(2-hydroxy-ethyl)-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl]-8-iodo-9H-purin-6-yl}-urea; 2-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-nicotinic acid; 5-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-pentanoic acid; 1-Ethyl-3-{9-[6-(2-hydroxy-propane-1-

sulfonylmethyl)-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl]-9H-purin-6-yl}-urea; 1-Ethyl-3-{9-[6-(2-methanesulfonylmethyl-pyrrolidine-1-carbonyl)-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl]-9H-purin-6-yl}-urea; 1-Ethyl-3-{9-[6-(2-methanesulfonylmethyl-pyrrolidine-1-carbonyl)-2-phenyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl]-9H-purin-6-yl}-urea; 1-Cyclopentyl-3-{9-[6-(2-methanesulfonylmethyl-pyrrolidine-1-carbonyl)-2-phenyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl]-9H-purin-6-yl}-urea; 6-[6-(3-Ethyl-ureido)-8-iodo-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxole-4-carboxylic acid; 3-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-2-hydroxy-propionic acid; and 3-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-2-oxo-propionic acid; or a pharmaceutically-acceptable salt, -hydrate, or -solvate thereof.

A particularly useful embodiment is where X in Formula II has a -CO₂H group on the ring.

In another embodiment, the compound is selected from the group consisting of: 3-{6-[6-(3-Ethyl-1-phenyl-ureido)-purin-9-yl]-2,2-dimethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-isoxazole-5-carboxylic acid; 3-{6-[6-(3-Ethyl-1-(5-methyl-furan-2-ylmethyl)-ureido)-purin-9-yl]-2,2-dimethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-isoxazole-5-carboxylic acid; 3-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-isoxazole-5-carboxylic acid; 3-{2,2-Dimethyl-6-[6-(3-phenyl-1-propyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-isoxazole-5-carboxylic acid; 5-Amino-2-{2,2-dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-N-hydroxy-benzamide; 6-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-nicotinamide; 1-{9-[6-(3-Hydroxy-pyridin-2-yloxymethyl)-2,2-dimethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl]-9H-purin-6-yl}-3-phenyl-urea; 3-{(2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxole-4-carbonyl}-amino)-benzoic acid; 2-{(2-Benzyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxole-4-carbonyl}-amino)-3-hydroxy-propionic acid; N-{2-Benzyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxole-4-carbonyl}-methanesulfonamide; 1-[9-(2-Benzyl-6-ureidomethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl)-9H-purin-6-yl]-3-phenyl-urea methylsulfonamide; 3-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-acrylic acid methyl ester; 3-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-propionic acid methyl ester; 3-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-

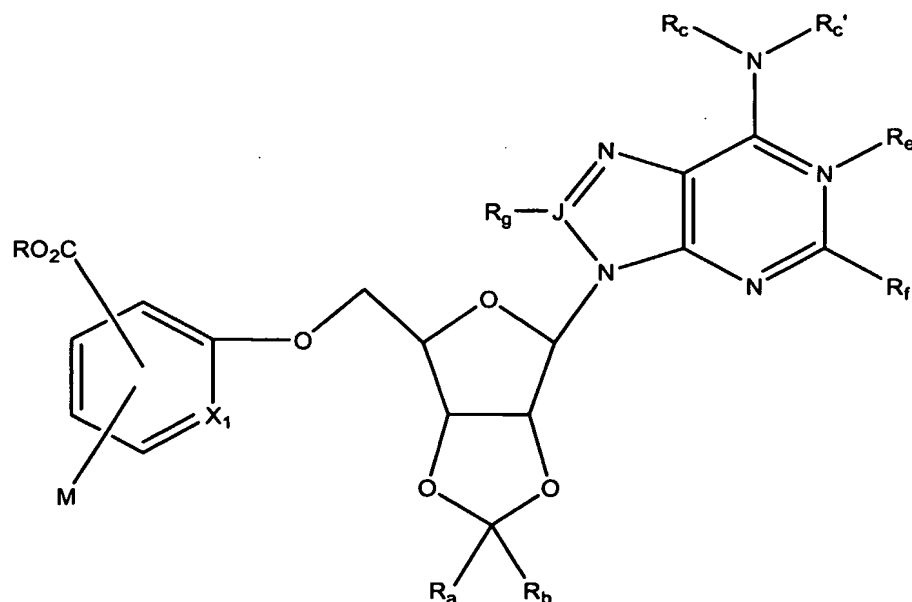
purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-yl}-propionic acid; and 3-(3-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-yl}-propionylamino)-benzoic acid; or a pharmaceutically-acceptable salt, -hydrate, or -solvate thereof.

In another embodiment of the method, R_b of Formula I is selected from the group consisting of: hydrogen, saturated or unsaturated C_{2-8} alkyl, saturated or unsaturated C_{3-7} cycloalkyl, aralkyl (including saturation and/or unsaturation in the alkyl portion), aryl, and C_{2-6} heterocycle; where all rings or chains optionally can bear one or more desired substituents.

In another embodiment of the method, X of Formula I is selected from the group consisting of: -OR, -SR, -NRR, -NR(CO)L, -NR(SO₂)L, -NR(CO)NR(SO₂)L, -NR(SO₂)NR(CO)L, and a moiety as provided in Formula II; Y, Z and Z' are independently selected from -CRR_r-, -O- and -S-; or the said -Y-Z'-, -(Z)₂-, -(Z)₃-, and -(Z)₄-units are selected from the respective said -Y-Z'-units and -(Z)_n-units as provided in Formula II.

In another embodiment of the method, compounds of general Formula I are molecules whose structures fall within the definition of Formula III:

Formula III



wherein R_a , R_b , R_c , R_c' , Σ , R , L , R_d , R_e , R_f , J , R_g are as defined in Formula I;

X_1 is selected from the group consisting of: N (nitrogen) and C-M; and

M is independently selected from the group consisting of: -H, halogen, CF₃, (saturated or unsaturated) C_{1-8} alkyl, (saturated or unsaturated) C_{3-7} cycloalkyl, aryl, aralkyl, heteroaryl,

(saturated or unsaturated) C₂₋₆ heterocycle, -OH, C₁₋₆ alkoxy, aralkoxy, aryloxy, -SH, C₁₋₆ thioalkyl, thioaryl, -[(CO)OR], -[(CO)NRR], amino, -N-substituted amino, and N,N-disubstituted amino; wherein each said substituent on said amino of M is independently selected from the group consisting of: (saturated or unsaturated) C₁₋₈ alkyl, (saturated or unsaturated) C₃₋₇ cycloalkyl, aryl, aralkyl, heteroaryl, (saturated or unsaturated) C₂₋₆ heterocycle, -[(CO)R], -[(CO)O-(C₁₋₈ alkyl)], and -[(CO)-NRR]; and when M is -[(CO)NRR], -[NH(CO)NRR], -[N(C₁₋₈ alkyl)(CO)NRR], -[N(aryl)(CO)NRR], or -[N(aralkyl)(CO)NRR], the R groups of any said -NRR unit (N,N-disubstituted-amino group) in M can be taken together such that a ring of 3 to 7 members is formed, with or without heteroatoms in place of the ring-carbon units; or a tautomer, or a pharmaceutically-acceptable salt, -hydrate, or -solvate thereof.

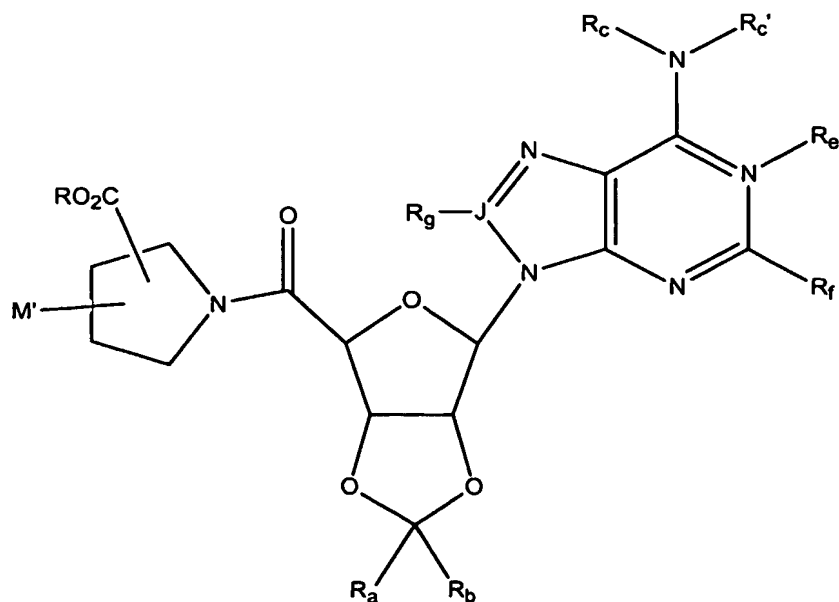
Another useful group of compounds are those of Formula III where the R group is H.

Examples of Formula III compounds are 5-Amino-2-{2,2-dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-benzoic acid; 4-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-isophthalic acid; 4-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-benzoic acid; 6-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-nicotinic acid; 5-Chloro-6-{2,2-dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-nicotinic acid; 2-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-nicotinic acid; 6-Chloro-2-{2,2-dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-5-fluoro-nicotinic acid; 6-Chloro-2-{2,2-dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-5-fluoro-nicotinic acid; 2-[6-[6-(3-Phenyl-ureido)-purin-9-yl]-2-(2-trifluoromethyl-phenyl)-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy]-nicotinic acid; 2-{2-Phenyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-nicotinic acid; 2-{2-Biphenyl-3-yl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-nicotinic acid; 2-{2-Naphthalen-2-yl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-nicotinic acid; 2-{2-Benzo[*b*]thiophen-3-yl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-nicotinic acid; 2-{6-[6-(3-Hexyl-ureido)-purin-9-yl]-2-phenyl-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-nicotinic acid; 2-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxo-spiroindan-4-ylmethoxy}-nicotinic acid; 2-{6-

[6-(3-Ethyl-ureido)-purin-9-yl]-2-phenethyl-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-
 nicotinic acid; 2-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-phenylethynyl-tetrahydro-furo[3,4-
d][1,3]dioxol-4-ylmethoxy}-nicotinic acid; 2-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-phenyl-
 tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-nicotinic acid; 2-{2-(2-Bromo-phenyl)-6-[6-
 5 (3-ethyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-nicotinic acid;
 2-{6-[6-(3-Cyclopentyl-ureido)-purin-9-yl]-2-phenethyl-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-
 ylmethoxy}-nicotinic acid; 2-{6-[6-(3-Cyclopentyl-ureido)-purin-9-yl]-2,2-(3,4-Dihydro-1H-
 naphthalen)-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-nicotinic acid; 2-{6-[6-(3-
 Cyclopentyl-ureido)-purin-9-yl]-2-*p*-tolyl-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-
 10 nicotinic acid; 2-{2-Biphenyl-4-yl-6-[6-(3-hexyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-
d][1,3]dioxol-4-ylmethoxy}-nicotinic acid; 2-{2-(4-Acetylamino-phenyl)-6-[6-(3-
 cyclopentyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-ylmethoxy}-nicotinic
 acid; and 2-{2-*tert*-Butyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-
d][1,3]dioxol-4-ylmethoxy}-nicotinic acid; or a pharmaceutically-acceptable salt, -hydrate, or
 15 -solvate thereof.

In another embodiment of the method, useful compounds fall within the definition of
 Formula IV:

Formula IV



wherein R_a , R_b , R_c , R_d , R_e , R_f , R_g are as defined in Formula I; and M' is as defined for M ;

the M' and $-CO_2R$ groups are independently attached to any carbon of the pyrrolidine ring; and

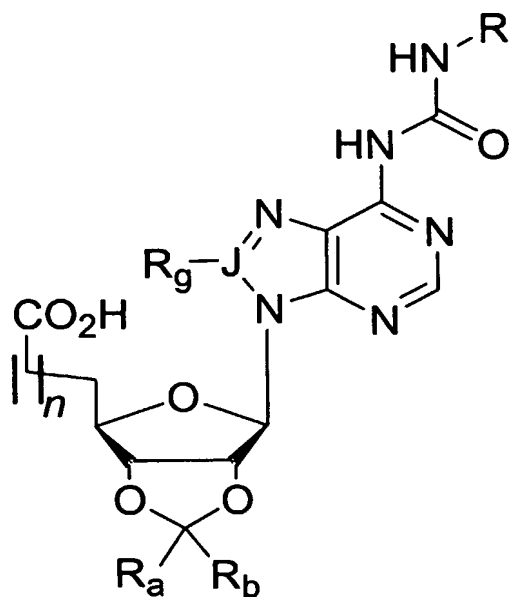
- 5 M' is not a halogen, hydroxy, sulfhydryl, or amino group when M' is attached to a carbon that is bonded to the pyrrolidine nitrogen atom (alpha position); or a tautomer, or a pharmaceutically-acceptable salt, -hydrate, or -solvate thereof.

Another useful group of compounds are those of Formula IV where $R = H$.

- Examples of Formula IV compounds are 1-{2-Phenyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxole-4-carbonyl}-pyrrolidine-2-carboxylic acid; 1-{2-Phenyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxole-4-carbonyl}-pyrrolidine-2-carboxylic acid; 1-{2-Benzyl-6-[6-(3-ethyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxole-4-carbonyl}-pyrrolidine-2-carboxylic acid; 1-(2-Phenyl-6-{6-[3-(2-phenyl-cyclopropyl)-ureido]-purin-9-yl}-tetrahydro-furo[3,4-*d*][1,3]dioxole-4-carbonyl)-pyrrolidine-2-carboxylic acid; 1-{6-[6-(3-Benzyl-ureido)-purin-9-yl]-2-phenyl-tetrahydro-furo[3,4-*d*][1,3]dioxole-4-carbonyl}-pyrrolidine-2-carboxylic acid; 1-{2-Benzo[*b*]thiophen-3-yl-6-[6-(3-hexyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxole-4-carbonyl}-pyrrolidine-2-carboxylic acid; 1-{2-Benzyl-6-[6-(3-hexyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxole-4-carbonyl}-pyrrolidine-2-carboxylic acid; 1-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-naphthalen-2-yl-tetrahydro-furo[3,4-*d*][1,3]dioxole-4-carbonyl}-pyrrolidine-2-carboxylic acid; 1-{6-[6-(3-Hexyl-ureido)-purin-9-yl]-2-phenyl-tetrahydro-furo[3,4-*d*][1,3]dioxole-4-carbonyl}-pyrrolidine-2-carboxylic acid; 1-{6-[6-(3-Cyclopentyl-ureido)-purin-9-yl]-2-phenyl-tetrahydro-furo[3,4-*d*][1,3]dioxole-4-carbonyl}-pyrrolidine-2-carboxylic acid; and 1-(3-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-*d*][1,3]dioxol-4-yl}-propionyl)-pyrrolidine-2-carboxylic acid; or a pharmaceutically-acceptable salt, -hydrate, or -solvate thereof.

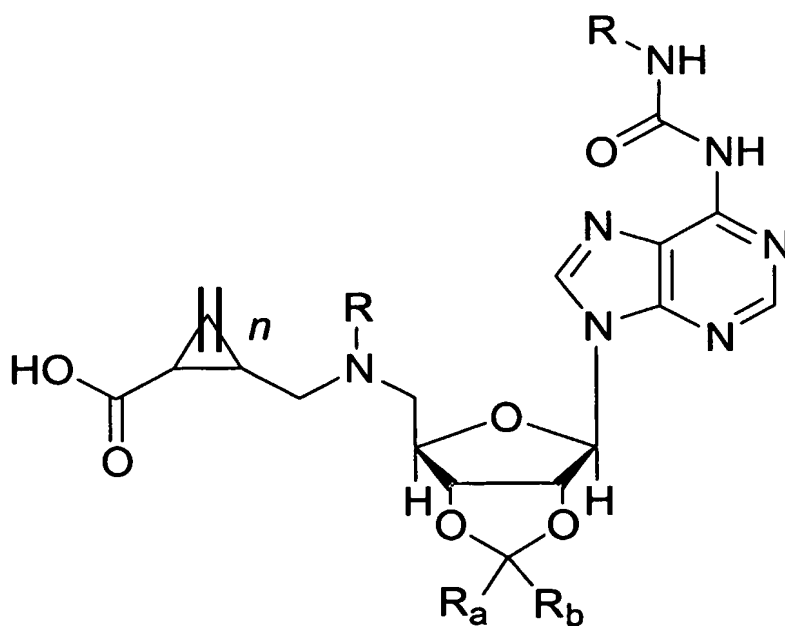
Other groups of compounds useful for the method are included under Formulas V-XI, or pharmaceutically acceptable salts, solvates, or hydrates thereof; in which R , R_a , R_b , J and R_g , are defined as for Formula I, and n is defined as for Formula II.

Formula V



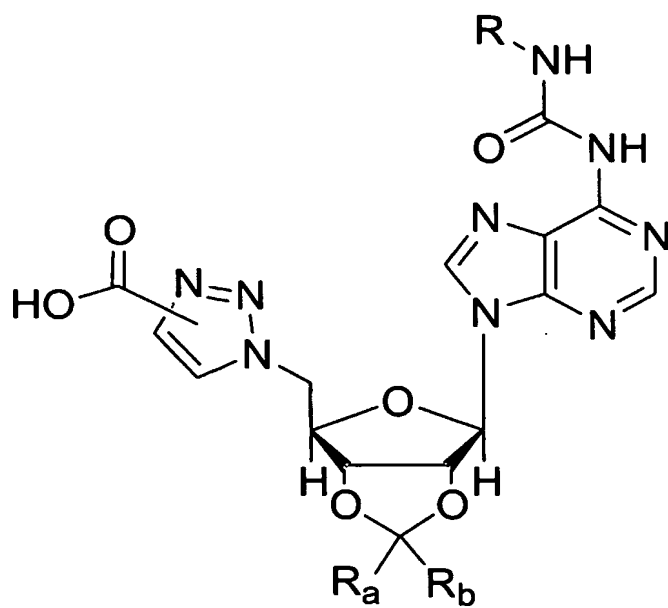
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Formula VI



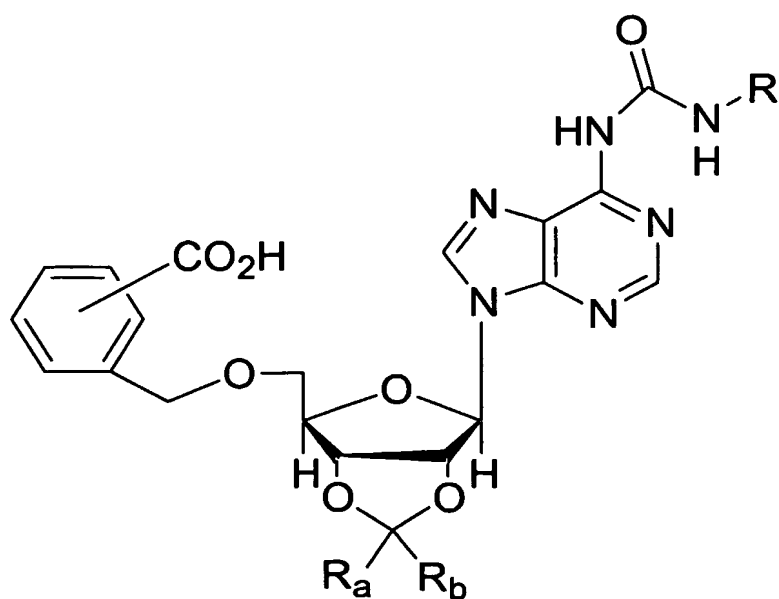
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Formula VII



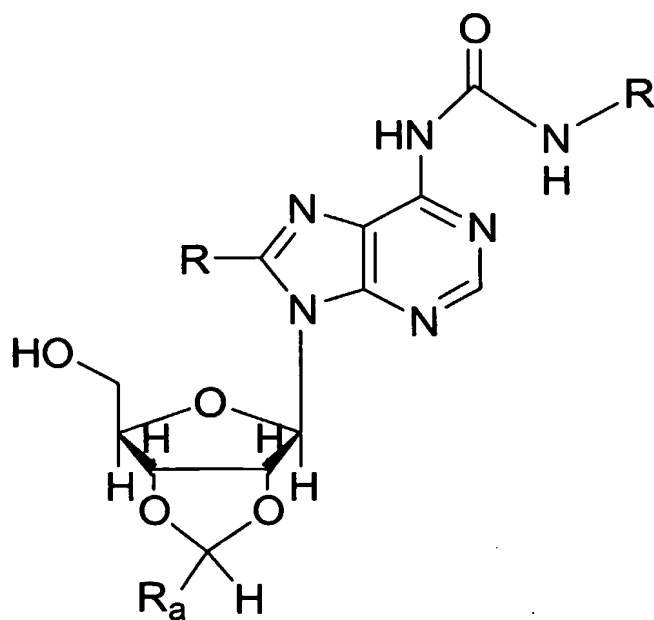
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Formula VIII



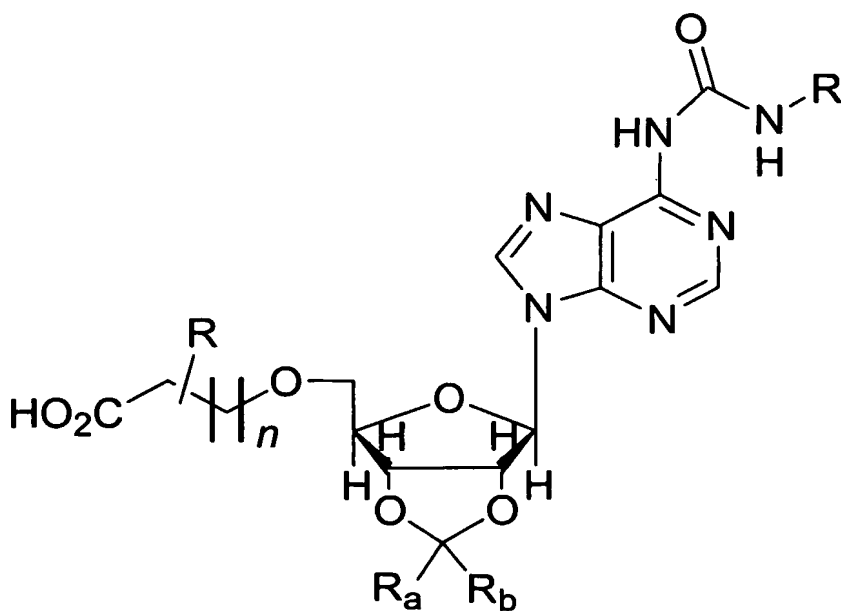
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Formula IX



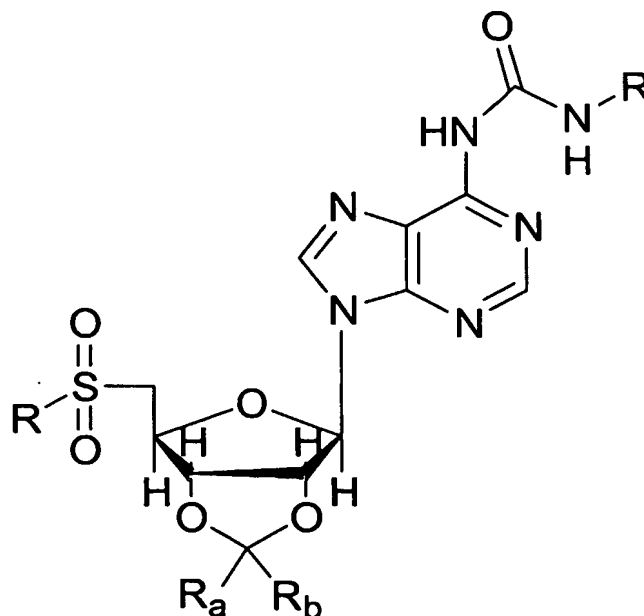
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Formula X



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Formula XI



5 Other groups of compounds useful for the method are included under Formulas V-XI, wherein the shown ureido-moiety in a given formula's structure (i.e., -N⁶HR_c of Formula I) is instead an amido-, guanidino-, or carbamoyl-moiety; or

wherein the shown ureido-moiety in a given formula's structure is not present, such that the compound is an N⁶-amine or N⁶-amine derivative; or

10 pharmaceutically acceptable salts, solvates, or hydrates thereof; in which R, R_a, R_b, J and R_g, are defined as for Formula I, and n is defined as for Formula II.

One embodiment of the present invention is a method to reduce pain through inhibition of P2X receptors with preferred compounds that are furanose-modified adenosine
15 analogues and/or their 8-aza-analogues.

Novel Compositions

The present invention also provides novel compounds of Formulas I, III, IV, V, VI, VII, VIII, IV, X, or XI, respectively, with the further provisos that for compounds of Formula

20 I:

(a) when R_a = Me, R_c = nothing, R_f = H, I or Cl, R_g = H, J = C, and the unit

-(R_c)N-(CG)-NR_dR_d = -HN-(CO)-NH-(aryl), -HN-(CO)-NH-(heteroaryl), -HN-(CO)-NH-(n-octyl), or -HN-(CO)-NH-(t-butyl), then R_b is not equal to Me; or

(b) when R_a = R_b = Me, R_e = nothing, R_f = R_g = H, J = C, and the unit R_c-N-(CG)-NR_dR_d' = (R')-N-(CO)-N(R'')(R'''), wherein R' is H, alkyl, substituted alkyl, heteroaryl-NH-(CG')-, or aryl-NH-(CG')-, G' is selected from the group consisting of: O, NR and S; R'' is H, alkyl, substituted alkyl, or aryl; and R''' is heteroaryl, aralkyl, or aryl, then the unit X-A-B- is not R-NH-(CO)-, unless said R-NH-(CO)- contains at least one -CO₂H group as a substituent on a ring or chain of said R; or

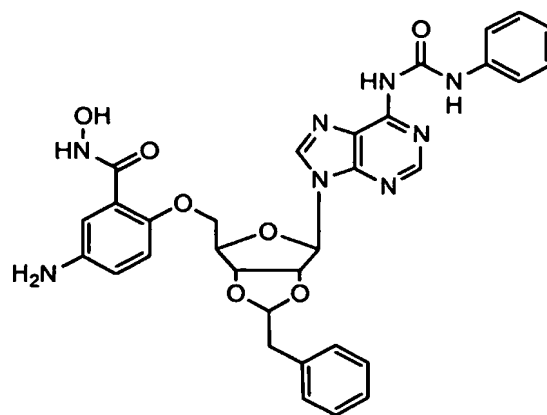
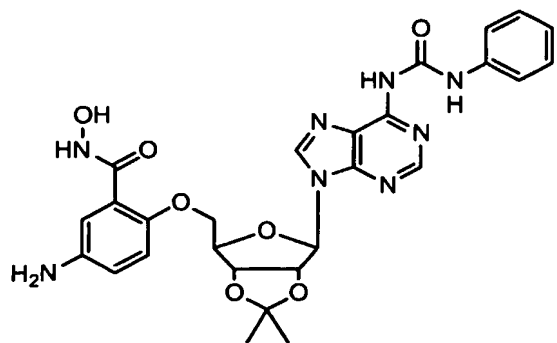
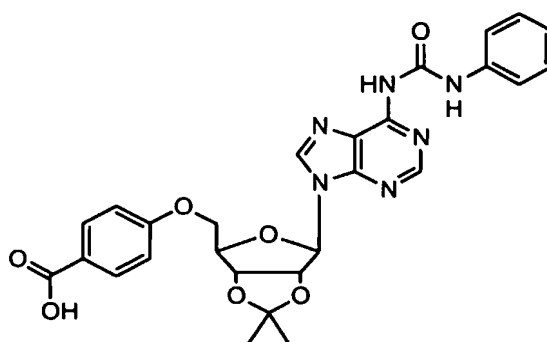
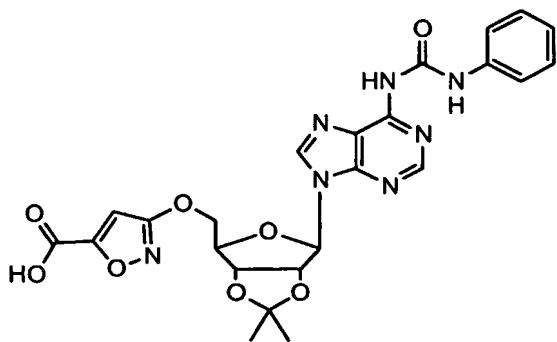
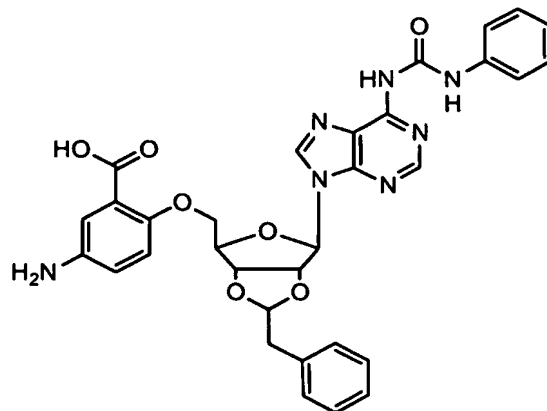
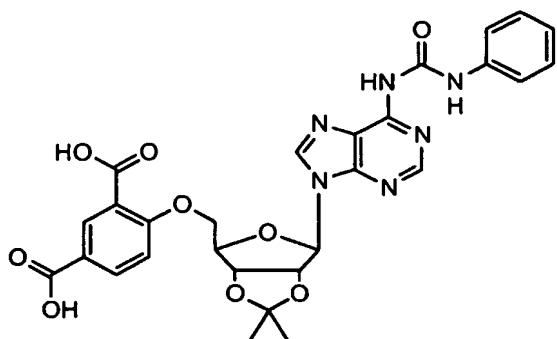
(c) when the unit X-A-B- = H(CO)-, RO(CO)-, ROCH₂-, RSCH₂-, RRNCH₂- then one of R_a or R_b must be H, and the other R_b or R_a is not H, Ph, Me or Et.

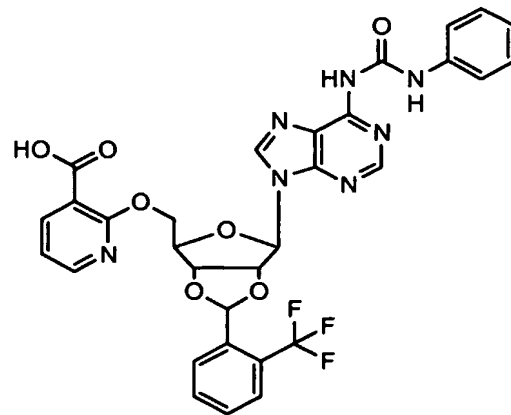
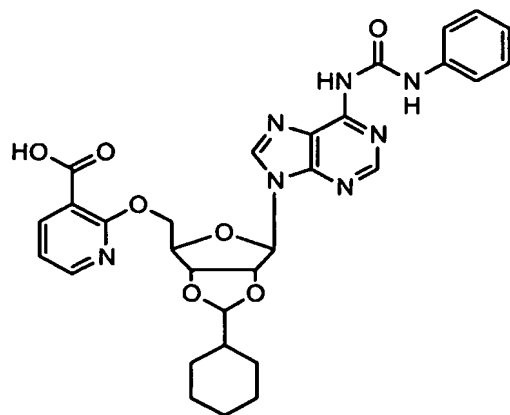
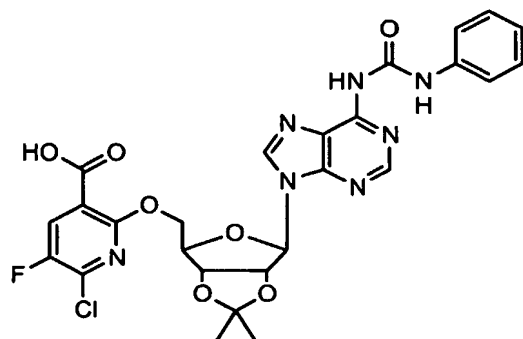
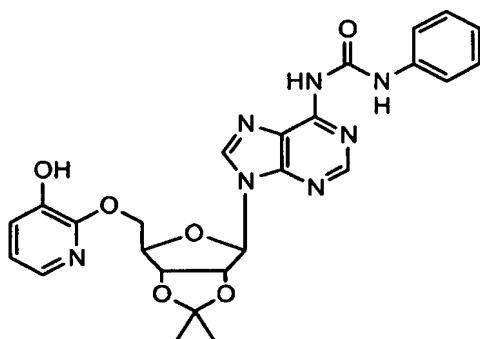
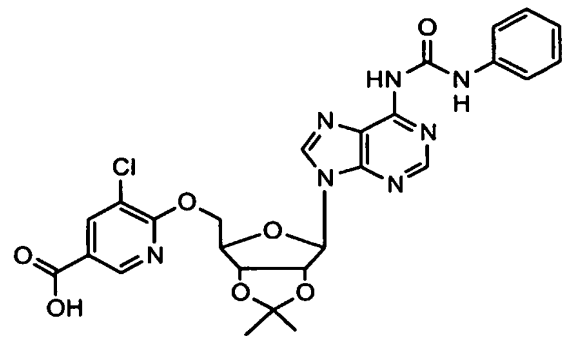
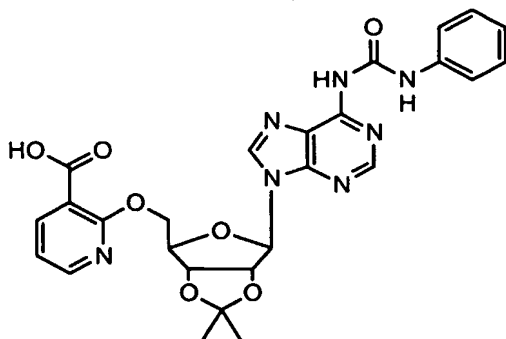
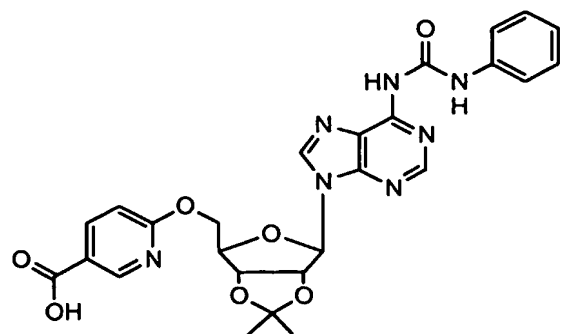
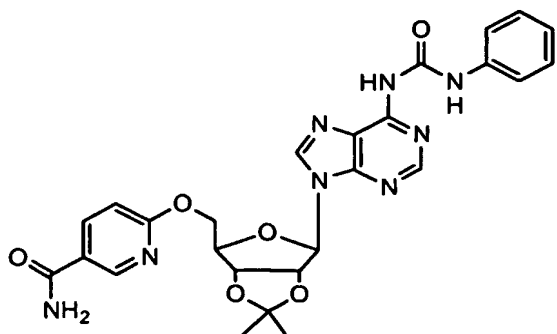
A further proviso for compounds of Formula V is: wherein at least one of R_a and R_b is not methyl.

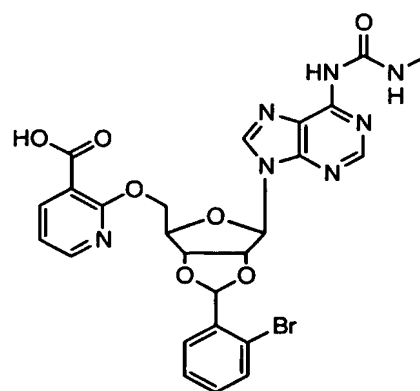
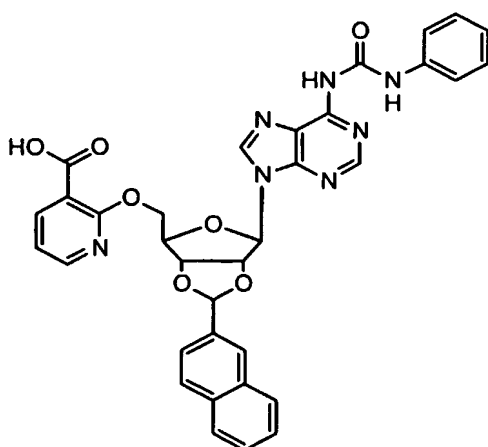
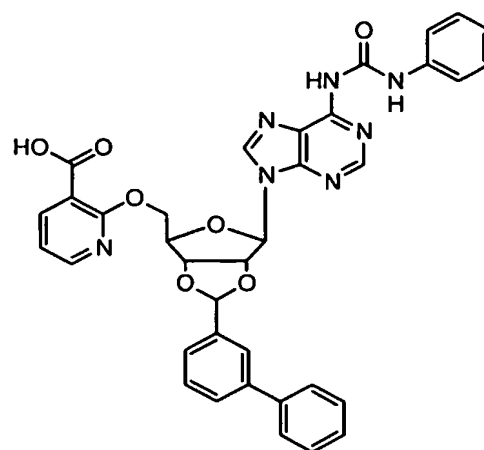
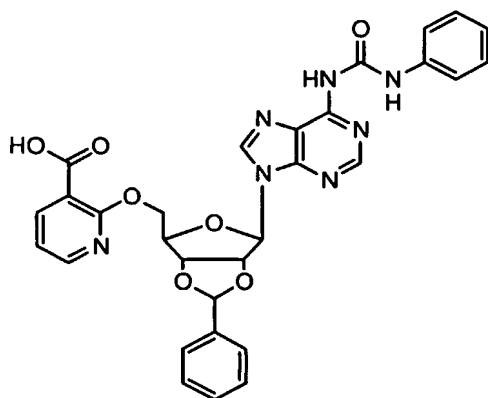
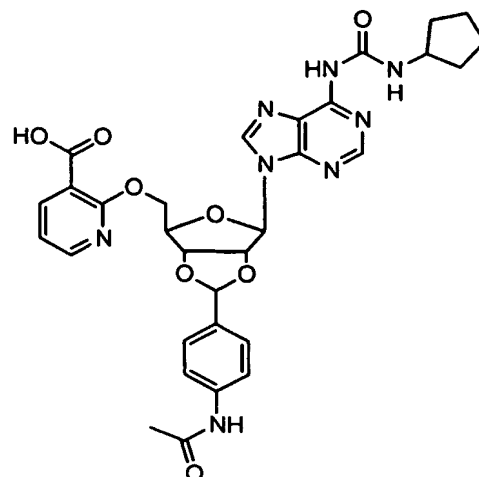
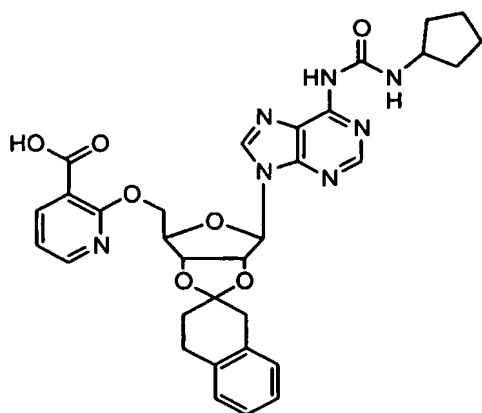
A desired substituent on a chain or ring (in place of a hydrogen at a position) is one selected from the given alkyl, aryl, halogen, aralkyl, carboxy, alkoxycarbonyl, hydroxyl, acyloxy, alkoxy, aryloxy or aralkoxy classes or from other classes, which provides a compound with good-to-excellent P2Y₁₂ receptor-binding properties, but which does not yield a compound with undesirable properties like chemical instability in a formulation, or one with levels of toxicity that are not well-tolerated by a treated mammal, or especially, not well-tolerated by a human.

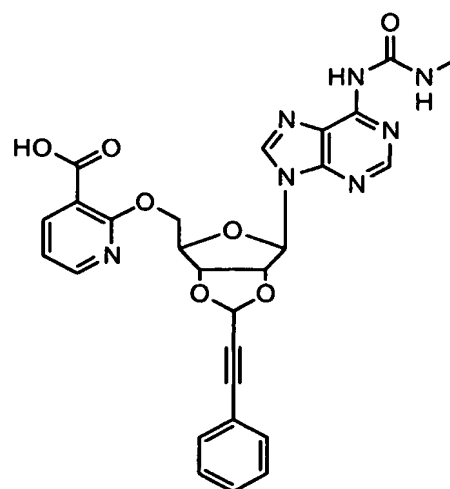
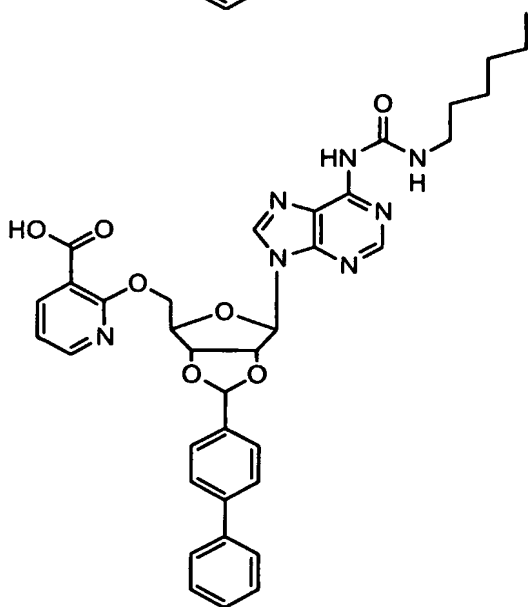
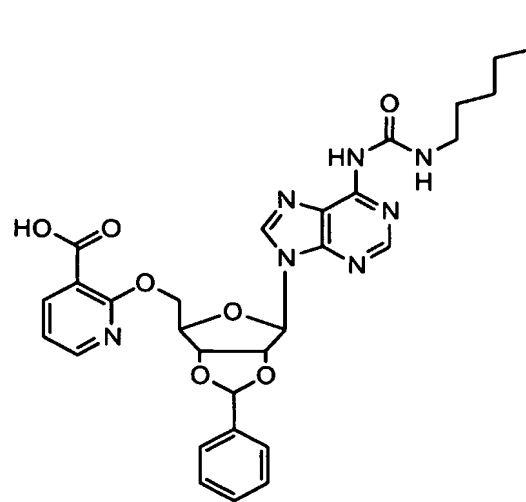
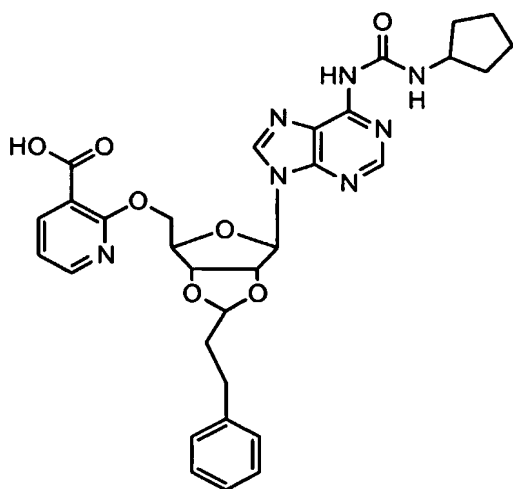
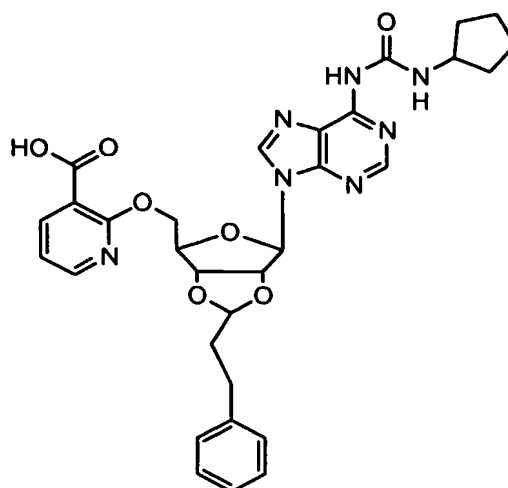
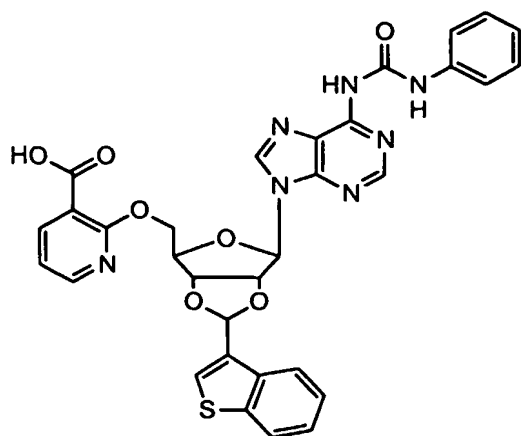
Exemplary compounds for the present invention are shown in Table 1.

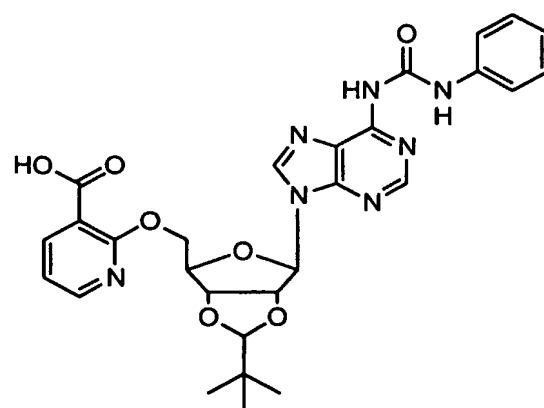
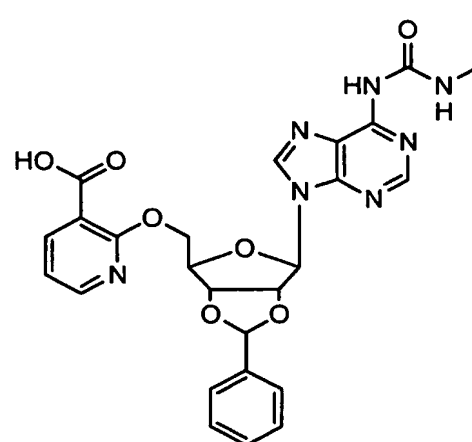
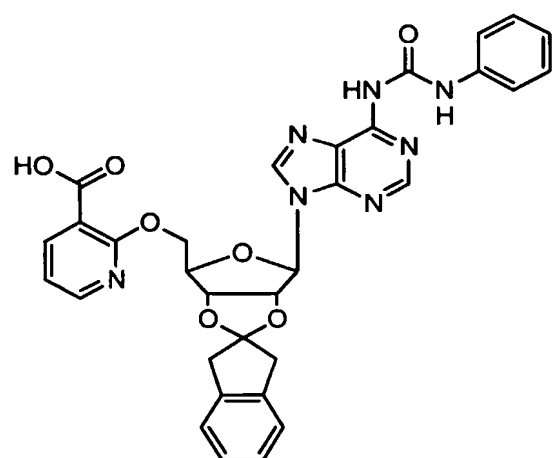
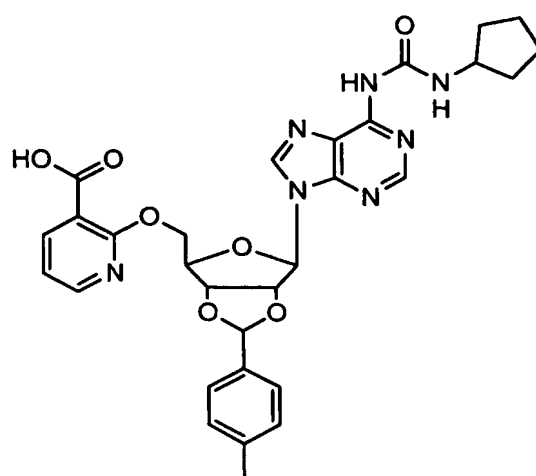
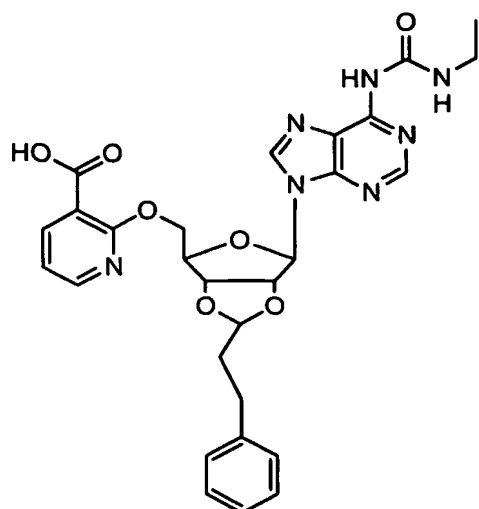
Table 1 - Exemplary Compounds.

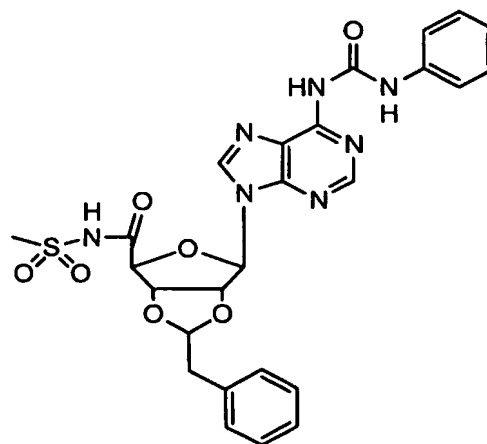
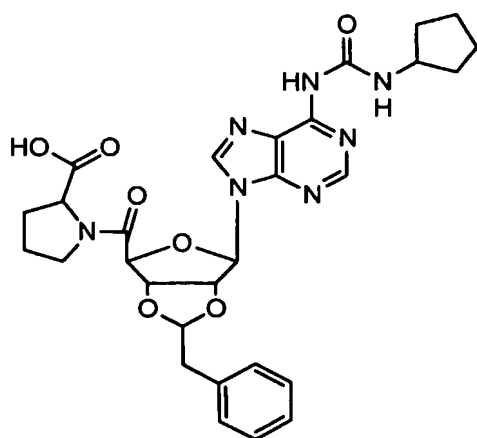
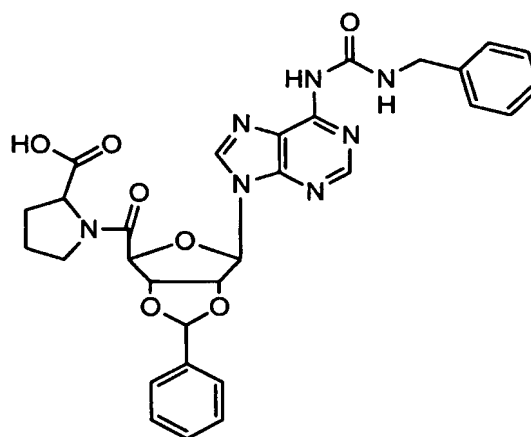
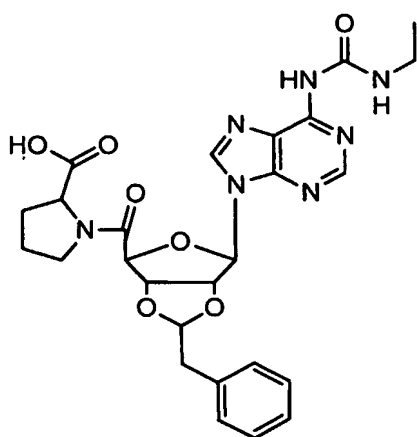
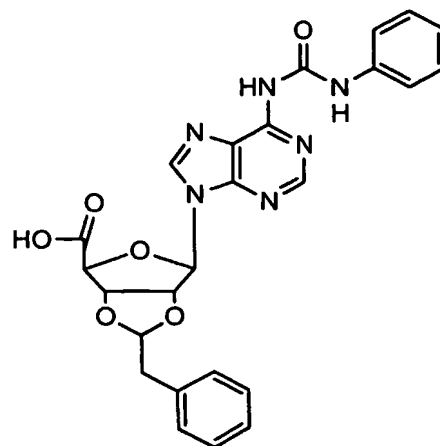
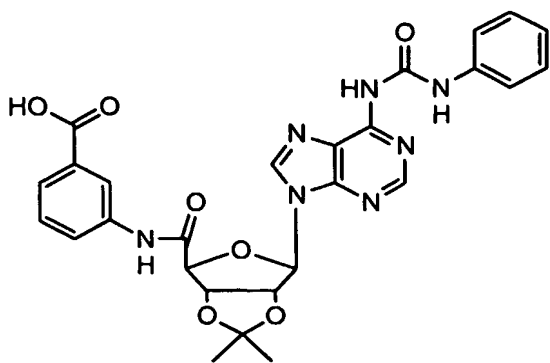


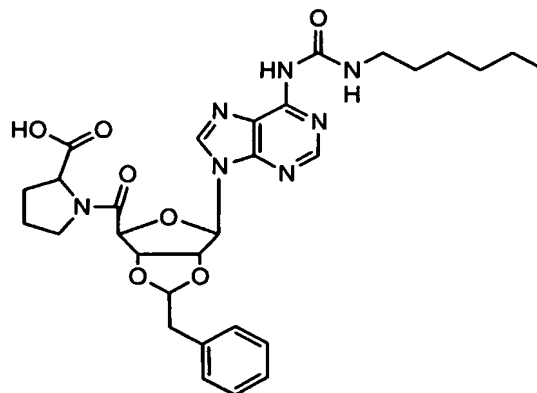
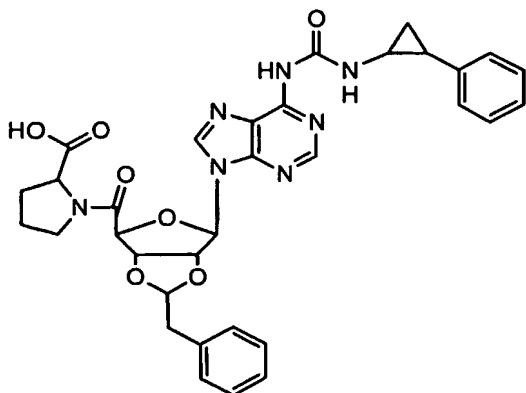
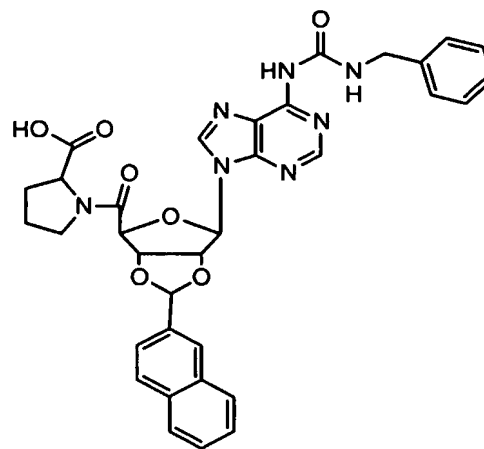
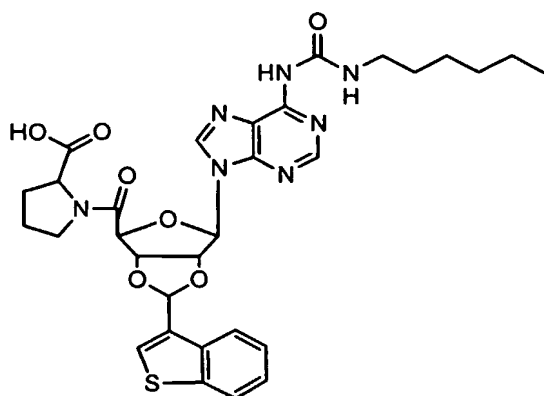
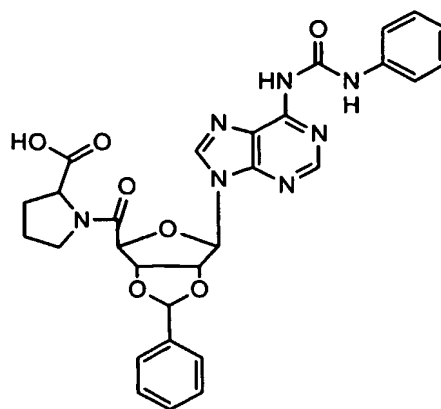
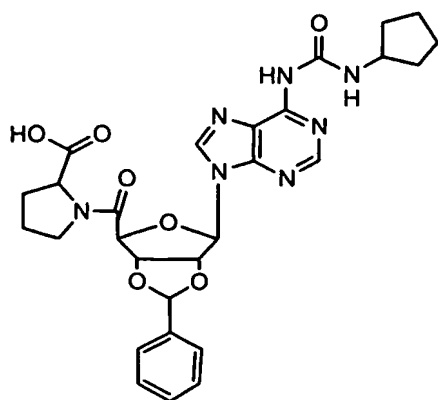


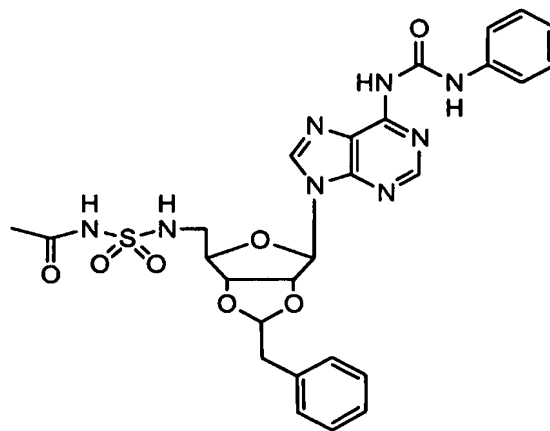
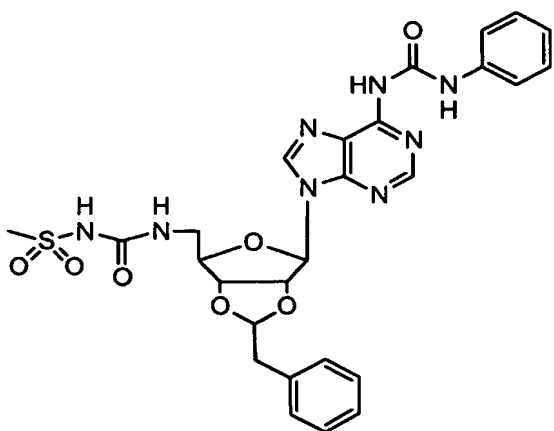
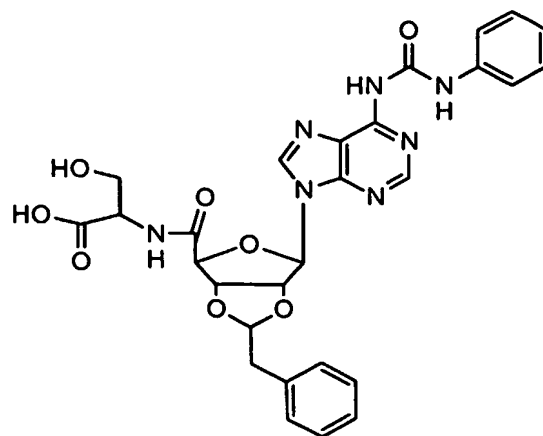
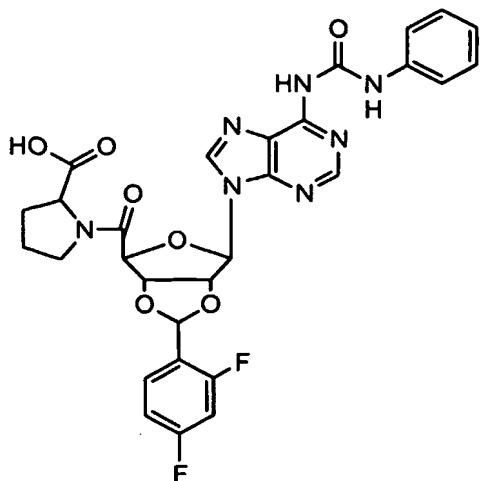


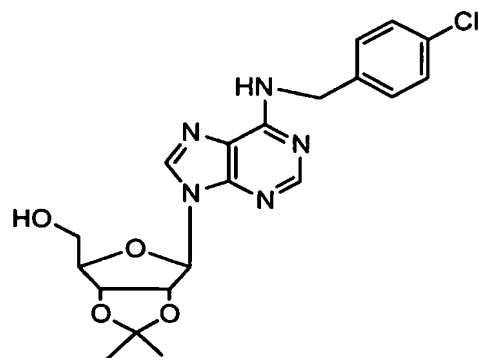
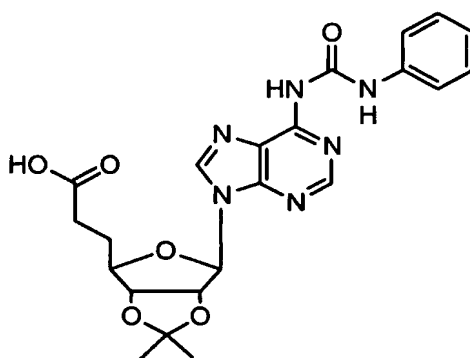
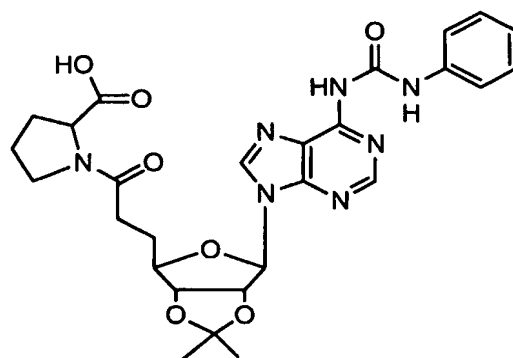
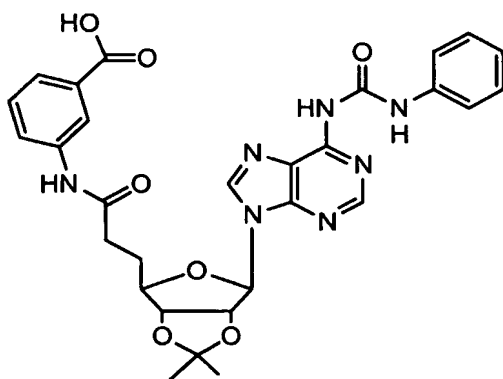
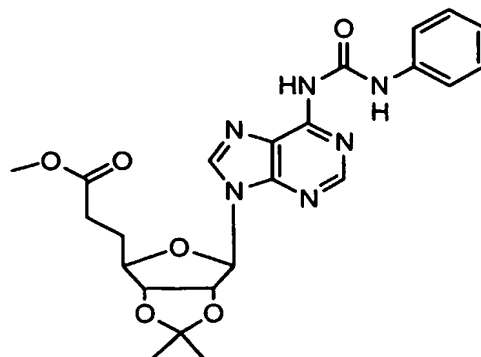
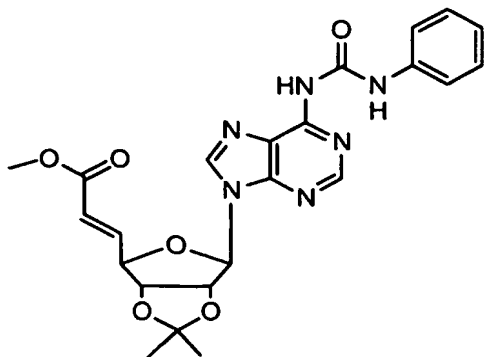


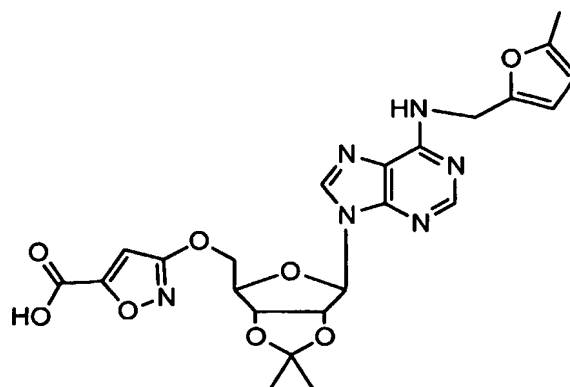
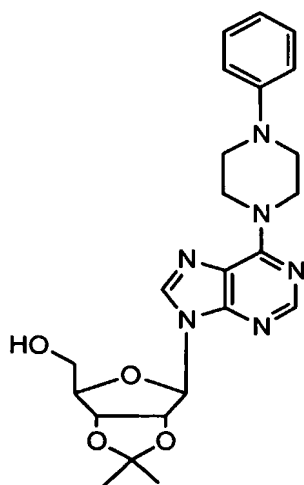
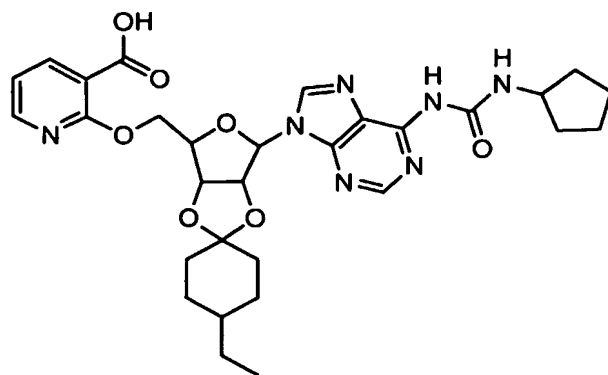
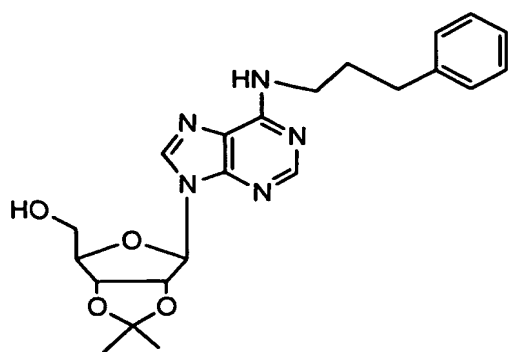


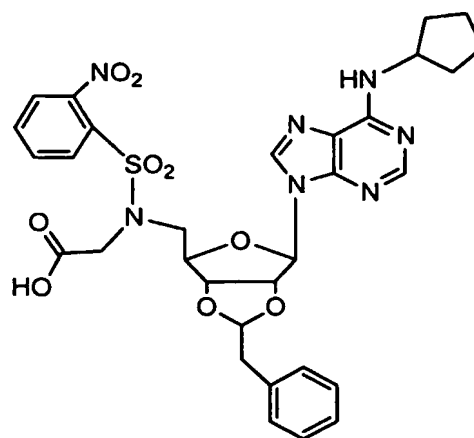
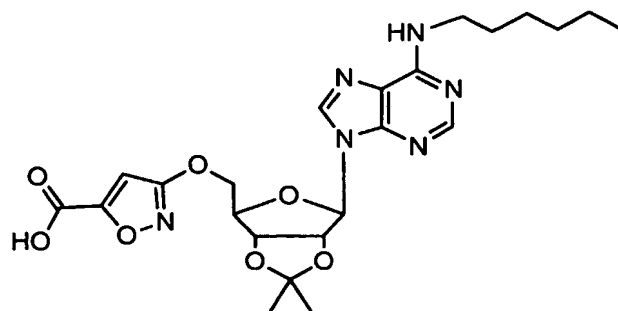
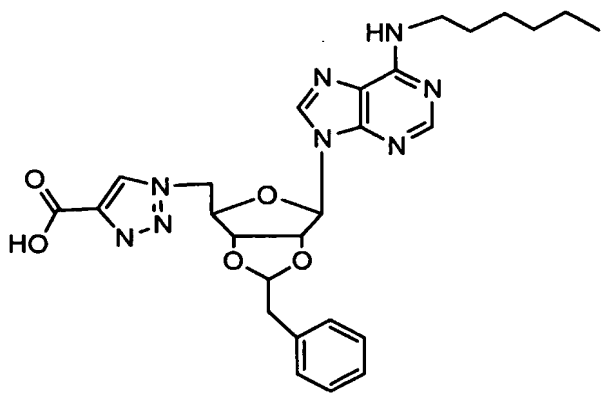
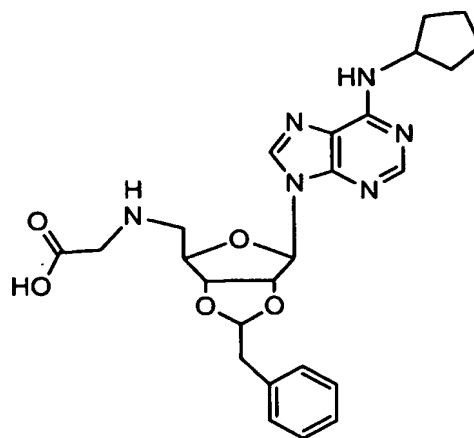
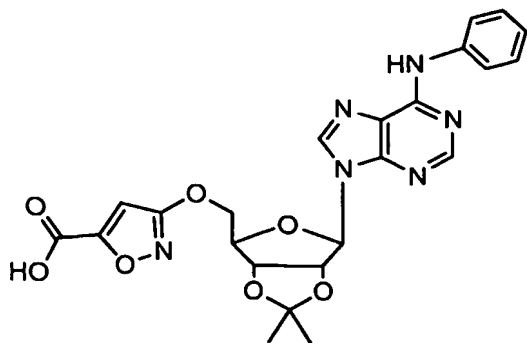












The present invention also encompasses non-toxic pharmaceutically acceptable salts
 5 of the above derivatives, such as, but not limited to, alkali metal salts such as lithium, sodium
 or potassium salts, or alkaline earth metal salts such as magnesium or calcium salts; or
 ammonium or mono-, di-, tri- or tetraalkyl ammonium salts, such as NH_4^+ , NEH_3^+ , NE_2H_2^+ ,
 NE_3H^+ , or NE_4^+ (wherein E is C_{1-4} alkyl) salts. Other salts such as hydrochlorides,

hydrobromides, mesylates, sulfates, acetates, tartrates, etc., are also contemplated in this invention. Pharmaceutically acceptable salts are salts that retain a desired biological activity level similar to that of the parent compound but that do not impart undesired toxicological effects. Preferred counterions are monovalent ions such as NH_4^+ , sodium, lithium, potassium, chloride, bromide, bisulfate, and mesylate, with sodium, potassium, chloride and mesylate being most preferred due to ease of manufacture, stability, and physiological tolerance.

Methods of Compound Preparation

The compounds of the present invention may be synthesized by those skilled in the art using conventional synthesis methodology and well-known workup and purification procedures. The following list of references, along with references cited therein, disclose general procedures employed for the synthesis of a number of intermediates and compounds related to the present invention: Baraldi, *et al.*, *Journal of Medicinal Chemistry*, 39(3): 802-806 (1996); Camaioni, *et al.*, *Bioorganic & Medicinal Chemistry*, 5(12): 2267-2275 (1997); Zablocki, *et al.*, PCT International Publication No. WO01/40243; Zablocki, *et al.*, PCT International Publication No. WO01/40246; Mantell, *et al.*, PCT International Publication No. WO01/94368; Jacobson, *et al.*, *Journal of Medicinal Chemistry*, 38(10): 1720-1735 (1995); Cristalli, *et al.*, *Journal of Medicinal Chemistry*, 38(9): 1462-1472 (1995); Secrist, III and Talekar, *Nucleosides & Nucleotides*, 9(4): 619-27 (1990); Secrist, III, USPN 4,794,174 (1988); Lyga and Secrist, III, *Journal of Organic Chemistry*, 48(12): 1982-1988 (1983); Dixon, *et al.*, PCT International Publication No. WO02/096248; Hardern, *et al.*, PCT International Publication No. WO01/36438; Guile *et al.*, PCT International Publication No. WO00/04021; Lee, *et al.*, *Bioorganic & Medicinal Chemistry Letters*, 13(6): 1087-1092 (2003); Cox, *et al.*, USPN 5,747,496 (1998).

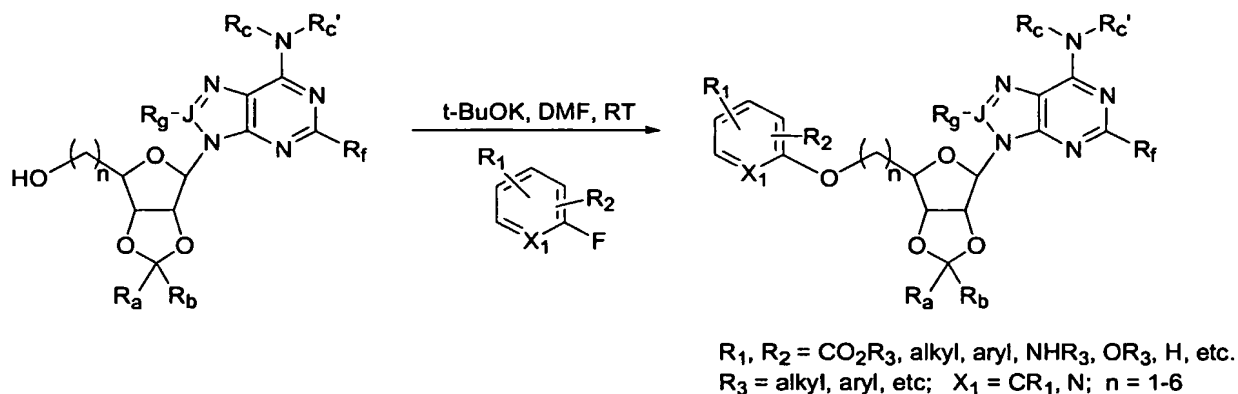
In many cases, commercially-available starting materials can be used for the synthesis of compounds of this invention. When not available commercially, useful starting materials can either be obtained from stepwise modification of commercially-available compounds and derivatives, or they may be synthesized from simpler precursors using literature methods known in the art. In addition, the compounds of the present invention may be synthesized using the general methods shown in Schemes 1-12, or variations thereof.

Commercially-available materials comprise: adenosine, α -adenosine, 2', 3'-isopropylideneadenosine, 5'-acetyl-2', 3'-isopropylideneadenosine, N^6 -(2-isopentenyl) adenosine, 2-chloroadenosine, 2-amino-6-chloropurine riboside, 6-chloropurine riboside,

inosine, 8-bromoguanosine, 8-bromoadenosine, 8-azidoadenosine, 8-azaguanine, 8-azaadenine, protected ribonic acid lactone derivatives and protected furanose derivatives. Other appropriate intermediates can be purchased from commercial sources and used as starting materials for compounds of the present invention, or can be synthesized as described in the chemical literature.

As disclosed above, commercially available compounds, or their derivatives may be employed as starting materials for the methods of Schemes 1-12.

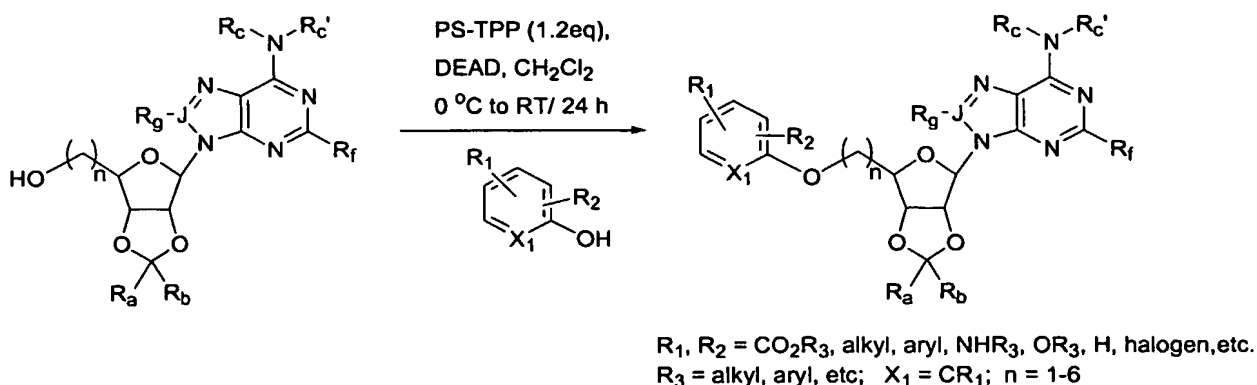
Scheme 1. Preparation of 5'-modified ethers by nucleophilic aromatic substitution.



Scheme 1, for example, discloses a useful method for the synthesis of 5'-aryl- or 5'-heteroaryl-nucleoside ethers and 8-azapurine-furanose 5'-ether derivatives by substitution of an appropriately functionalized adenosine analogue or 8-azapurine derivative for a halogen on an appropriately substituted halogenated aromatic compound or a related heteroaromatic derivative. Generic groups in Scheme 1 are as described in Formula I, which also includes the general definitions at the end of this section; some examples of preferred groups for R_1 , R_2 , and R_3 are provided, but these examples should not be considered limiting. For example, substituents at R_1 and R_2 of the aromatic/heteroaromatic-group in Scheme 1 can independently be hydrogen, or alkyl, or groups containing carboxylic acid derivatives such as: $-\text{CO}_2R_3$; but they can also be halogen, or esters or amides of alkylcarboxylic acids, arylcarboxylic acids, $-\text{O}-(\text{alkylcarboxylic acids})$, $-\text{NR}-(\text{alkylcarboxylic acids})$, and the like. When either or both of R_1 and R_2 are halogen in Scheme 1, preferred halogens are chloro and fluoro. It will be understood by those skilled in the art that, in some cases, the N^6 -group will

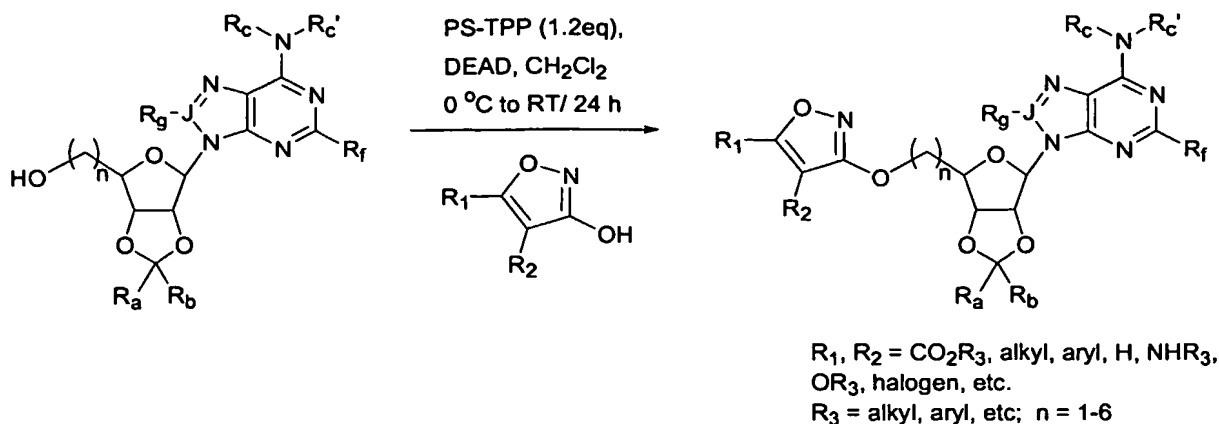
have to be temporarily protected using a group or groups such as benzyl, phthaloyl, etc., in order to allow the desired coupling in Scheme 1 to occur without competition by the N⁶-group. Removal of said protecting group using methods known in the art then yields a compound of the present invention. In many cases, chemical intermediates produced using methods outlined in Schemes 1-12 may be useful in their own right as compounds useful for the practice of the methods of this invention.

Scheme 2. Preparation of 5'-modified ethers by Mitsunobu coupling of phenols.



5'-Substituted aryl nucleoside derivatives can also be prepared via Mitsunobu coupling (Mitsunobu, *Synthesis* 1-28 (1981); Brown, et al., *J. Med. Chem.* 37 (5), 674-88 (1994); Santosh and Balasubramanian, *Synthetic Communications*, 24(8), 1049-62 (1994)) of phenols to derivatives of adenosine, 8-azaadeonsine, guanosine, 8-azaguanosine, etc., as provided in Scheme 2. Generic groups in Scheme 2 are as described in Formula I, which also includes the general definitions at the end of this section; some examples of preferred groups for R₁, R₂, and R₃ are provided, but these examples should not be considered limiting. For example, substituents at R₁ and R₂ of the aromatic/heteroaromatic-group in Scheme 2 independently can be hydrogen, halogen, alkyl, alkoxy, aryl or groups containing carboxylic acids, and/or their derivatives such as: -CO₂R₃; but esters or amides of alkylcarboxylic acids, arylcarboxylic acids, -O-(alkylcarboxylic acids), -NR-(alkylcarboxylic acids), and the like are also included. When either or both of R₁ and R₂ are halogen in Scheme 2, preferred halogens are chloro and fluoro.

Scheme 3. Preparation of 5'-modified isoxazole ethers by Mitsunobu coupling of isoxazole derivatives.



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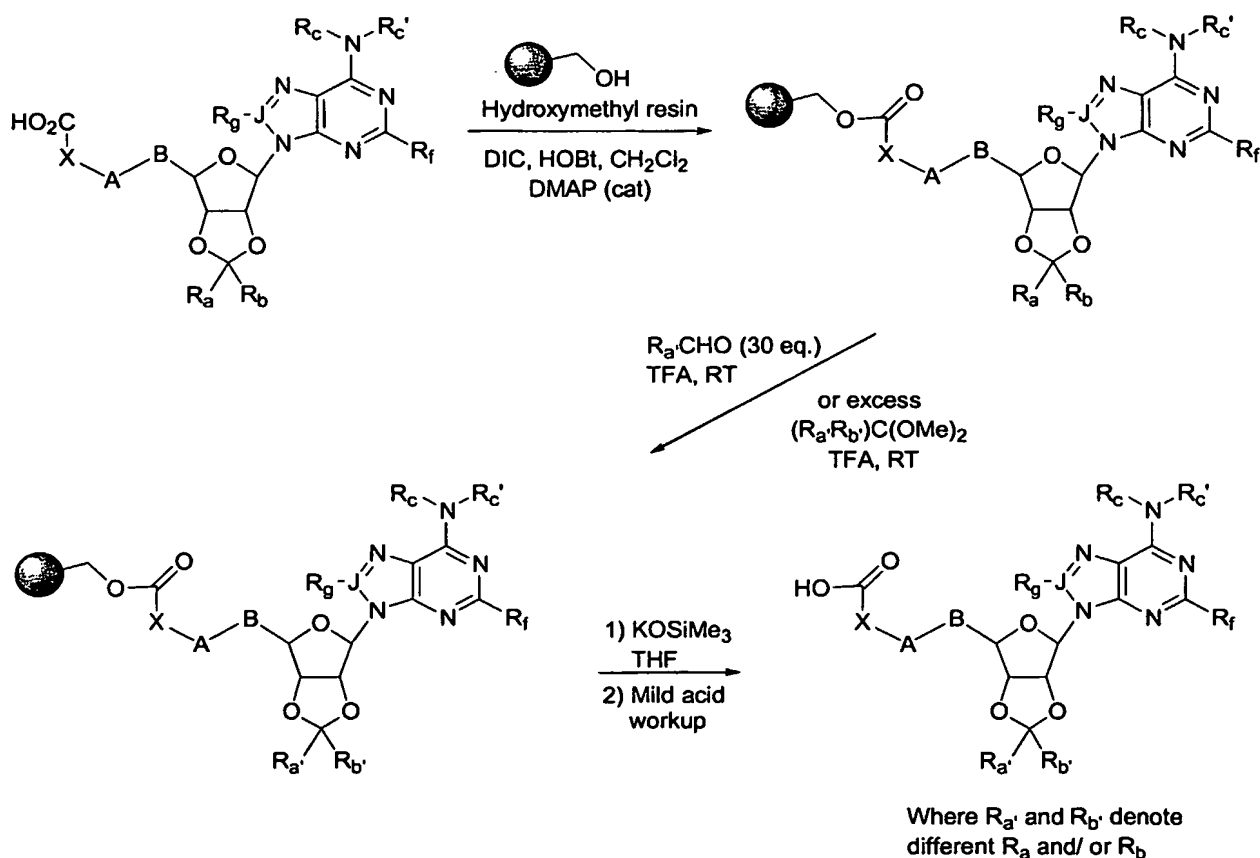
Alternatively, the Mitsunobu coupling may be carried out using hydroxyisoxazoles and the appropriate derivatives of adenosine, 8-azaadenosine, guanosine, 8-azaguanosine, etc., as provided in Scheme 3. Generic groups in Scheme 3 are as described in Formula I, which also includes the general definitions at the end of this section; some examples of preferred groups for R_1 , R_2 , and R_3 , are provided in the scheme, but these examples should not be considered limiting. For example, preferred substituents at R_1 and R_2 of the isoxazole derivatives of Scheme 3 independently are hydrogen, alkoxy, or halogen, or groups containing carboxylic acids or their derivatives such as: $-\text{CO}_2R_3$; but esters or amides of alkylcarboxylic acids, arylcarboxylic acids, $-\text{O}-(\text{alkylcarboxylic acids})$, $-\text{NR}-(\text{alkylcarboxylic acids})$, and the like, are also included. When either or both of R_1 and R_2 are halogen in Scheme 3, preferred halogens are chloro and fluoro.

When the products of any of Schemes 1, 2 or 3 comprise esters, said esters themselves can be useful in the invention. Said ester derivatives may be purified by methodologies well-known in the art, such as by normal phase, or reverse phase chromatography, or in suitable circumstances, using crystallization techniques. Alternatively, said ester derivatives can be used in the synthesis of other derivatives such as amides, hydroxamic acids, and different alkyl esters by well-known methods in the art.

Optionally, said esters may be hydrolysed under basic conditions, or cleaved using other methods known in the art, which cleave esters selectively in the presence of ketals or acetals to provide acid salts. These salts are also useful in the invention.

If desired, said acid salts can be converted into acids upon mild acid treatment. Workup by common techniques and purification by methods well known in the art, including purification by crystallization or chromatography, can be used to give the purified acids. Said acids can also be converted into other useful derivatives such as amides, hydroxamic acids, aryl esters, etc., by methods known to those skilled in the art of chemical synthesis. These acid derivatives are also useful in the invention, and are also purified using well-known methods such as crystallization or chromatography.

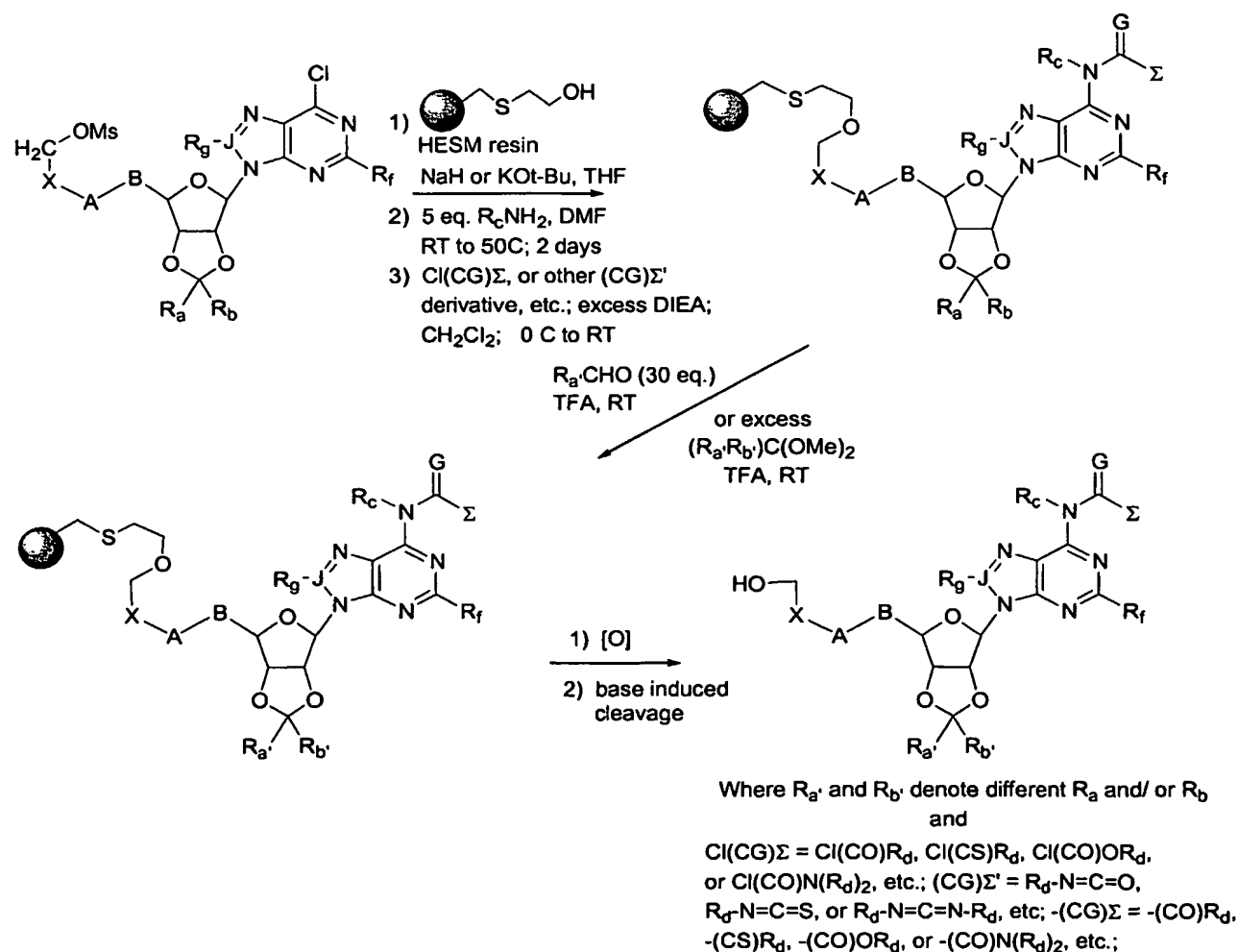
Scheme 4. Transformation of 2', 3'-acetals and -ketals into new acetal and ketal derivatives using solid phase techniques.



Diversity using common intermediates can be introduced into the 2', 3'-acetal or -ketal position of compounds encompassed in Formula I, as well as at the N⁶-position of compounds of Formula I using solid-phase synthetic methods. Schemes 4 and 5 exemplify transketalization procedures using polymer-bound relatives of compounds of Formula I. These methods can be used to transform one class of 2', 3'-ketal or -acetal into other useful

2', 3'-acetal or -ketal derivatives of adenosine, guanosine, 8-azaadenosine, etc. These polymer-bound approaches are very useful for after the desired reaction is complete, excess reagents can be washed away using one to several solvent washes. The desired material remains attached to the resin, in a pre-purified form, until it is cleaved from the solid phase using the appropriate conditions. Final purification of the desired product is then accomplished by conventional techniques like chromatography or crystallization.

Scheme 5. Transformation of 6-chloro derivatives into 6-amino, 6-amido, 6-carbamoyl, 6-ureido, 6-thioureido, 6-guanidino, etc., derivatives and subsequent conversion of 2', 3'-acetals and -ketals into new acetal and ketal derivatives using solid phase techniques.



Scheme 5 shows another variation of compound preparation employing resin-bound materials of Formula I. It outlines a sequence of solid-phase procedures that are useful for the

introduction of functionality at the 6-position, as well as for transketalization at the 2', 3'-position, if desired. It should be noted that the chemistry procedures used in these solid phase approaches are similar to methods known and used for solution-phase chemical transformations involving the synthesis of adenosine, guanosine, 8-azaadenosine, etc., derivatives. The primary differences are the necessity of attachment of a starting material to a resin, the simplicity of resin-based purification techniques (filtration and washing by compatible solvents) compared to solution-phase purification techniques (chromatography crystallization, etc.), and the requirement for cleavage of a compound of the invention, or an intermediate useful for the synthesis of a compound of the invention, from a resin prior to final purification and/or use of a compound so cleaved.

In Scheme 5, an early intermediate, such as a 6-chloroadenosine-2', 3'-ketal or -acetal derivative, or a 6-chloro-8-aza-adenosine-2', 3'-ketal or -acetal derivative is attached to a resin such as polystyrene resin via a β -thioethanol linker (e.g., hydroxyethylsulfanylmethyl polystyrene; HESM polystyrene resin; Garcia-Echeverria, *Tetrahedron Lett.*, 38, 8933-7 (1997)). After workup by filtration and rinsing with a useful solvent like DMF, the resin bound material is treated with a primary or secondary amine, ammonia or a hydroxylamine derivative to introduce an amino group via displacement of the 6-chloride. Scheme 5 exemplifies the process using a generic primary amine, R_cNH_2 . Though not shown in Scheme 5, a resin-bound product from chloride displacement by R_cNH_2 can be cleaved from the solid-phase after formation and workup without conversion into an amide or other derivative; such an amine product, in itself, can be useful for this invention. Or, the said resin-bound C⁶-amine product, itself, can be used in the transketalization/transacetalization step shown in Scheme 5 without prior transformation into a -(CG) Σ derivative, or a -(CG) Σ' derivative, to give a different 2', 3'-acetal-6-amine derivative tethered to the solid phase. When cleaved from the solid phase, such a C⁶-amine material can also be useful in the method of the invention. If desired, such a C⁶-amine material can be further converted into one or more other N⁶-derivatives as shown. It should be noted that treatment of a 6-chloro derivative, such as shown in Scheme 5, with secondary amines, such as $(R_c)_2NH$ or $R_cR_{c'}NH$, can also yield products useful in this invention. For example, an embodiment of this invention comprises compounds that have a C⁶-piperazino-group; a preferred embodiment of this class of compounds comprises a substituted piperazino-group on C⁶. A particularly useful class of derivatives comprises an aryl substituent on the distal amine of said piperazino group. Alternatively, if a (R_c) -group is used as a temporary protecting group,

such as for the selective introduction of a N⁶-group or for protection during a later step, said (R_c)-group can be removed after it has served its purpose in a desired chemical step or steps to yield a compound useful in the invention, or a compound useful for conversion into a such a compound of the invention.

5 As with solution phase techniques, subsequent introduction of an amido-, thioamido-, carbamoyl-, ureido-, thioureido-, or guanidino-group at N⁶ of a polymer-bound amine intermediate such as provided for in Scheme 5 can be made in one step using an excess of an appropriate acid/ coupling agent combination in an appropriate solvent (e.g., methoxypropionic acid/dicyclohexylcarbodiimide/ dichloromethane), or using other
10 conditions known in the art, such as those involving an acid chloride, acid anhydride, carbonyl chloride, chloroformate, isocyanate, isothiocyanate, carbodiimide, carbamoyl chloride, or 2-alkyl-2-thiopseudourea; or using a chemical equivalent of such materials using well-known methods in the art, or using other methods and reagents in the literature.

Alternatively, a two-step approach can be used to introduce a group at N⁶, which
15 comprises treatment of an appropriate 6-amino derivative, synthesized as in Scheme 5, with a solution of a small excess of phosgene or thiophosgene and a tertiary amine such as DIEA in a suitable solvent such as dichloromethane or toluene at a temperature which allows reaction, followed by treatment with an excess of a nucleophile, such as a primary or secondary amine to give a resin-bound urea or thiourea after workup. Other useful nucleophiles for this
20 reaction are alcohols, phenols, hydrazine derivatives, etc. If desired, transketalization can be performed on a bound substrate of Scheme 5 in a manner similar to that shown in Scheme 4. Or, if a desired acetal or ketal moiety of a 6-chloroadenosine, 6-chloroguanosine, 6-chloro-8-azaadenosine, etc., derivative is used to begin with, then no transketalization is necessary.

Cleavage from the β-thioethanol linker of the solid phase, as shown for the HESM
25 polystyrene resin in Scheme 5 is performed in two steps: oxidation of the thioether-linker using an oxidizing agent like MCPBA in a solvent such as dichloromethane gives a sulfone-linker, then cleavage of the β-ether moiety from the oxidized linker occurs upon treatment with a strong base like DBU in a solvent like dichloromethane and yields a compound which can be purified by techniques known in the art. Preferred oxidizing agents include peracids
30 like MCPBA and peracetic acid, but other oxidizing agents like hydrogen peroxide, permanganate salts, or persulfate salts can be used to oxidize a thioether to a sulfone. Preferred elimination conditions include DBU in dichloromethane, or 10% ammonium hydroxide in trifluoroethanol.

Following cleavage from the solid phase, a compound formed using the procedures in Scheme 5 can be used in the present invention, or can be further modified by well-known methods for functional-group transformations to generate new compounds which are also useful for this invention.

5 Preferred aldehydes, aldehyde acetals and ketone ketals useful in the transketalization methods shown in Schemes 4 and 5 comprise the below said carbonyl compounds and/or derivatives of: benzaldehyde, biphenyl-3-carboxaldehyde, biphenyl-4-carboxaldehyde, biphenyl-4-yl-acetaldehyde, 2-bromobenzaldehyde, benzo[b]thiophene-3-carbaldehyde, cyclohexanecarbaldehyde, cyclopentanecarbaldehyde, 2,5-dimethylbenzaldehyde, 2,6-
10 difluorobenzaldehyde, 2-fluorobenzaldehyde, naphthalene-2-carbaldehyde, phenyl acetaldehyde, phenyl propynal, 3-phenyl propenal, 3-phenyl propionaldehyde, 2-trifluoromethyl benzaldehyde, cyclohexanone, cyclopentanone, 4-ethyl cyclohexanone, 3,4-dihydro-1H-naphthalen-2-one, indan-2-one, and propan-2-one. Useful derivatives for transketalization of the above ketones comprise ketals like dimethoxy- or diethoxy-ketals, etc.

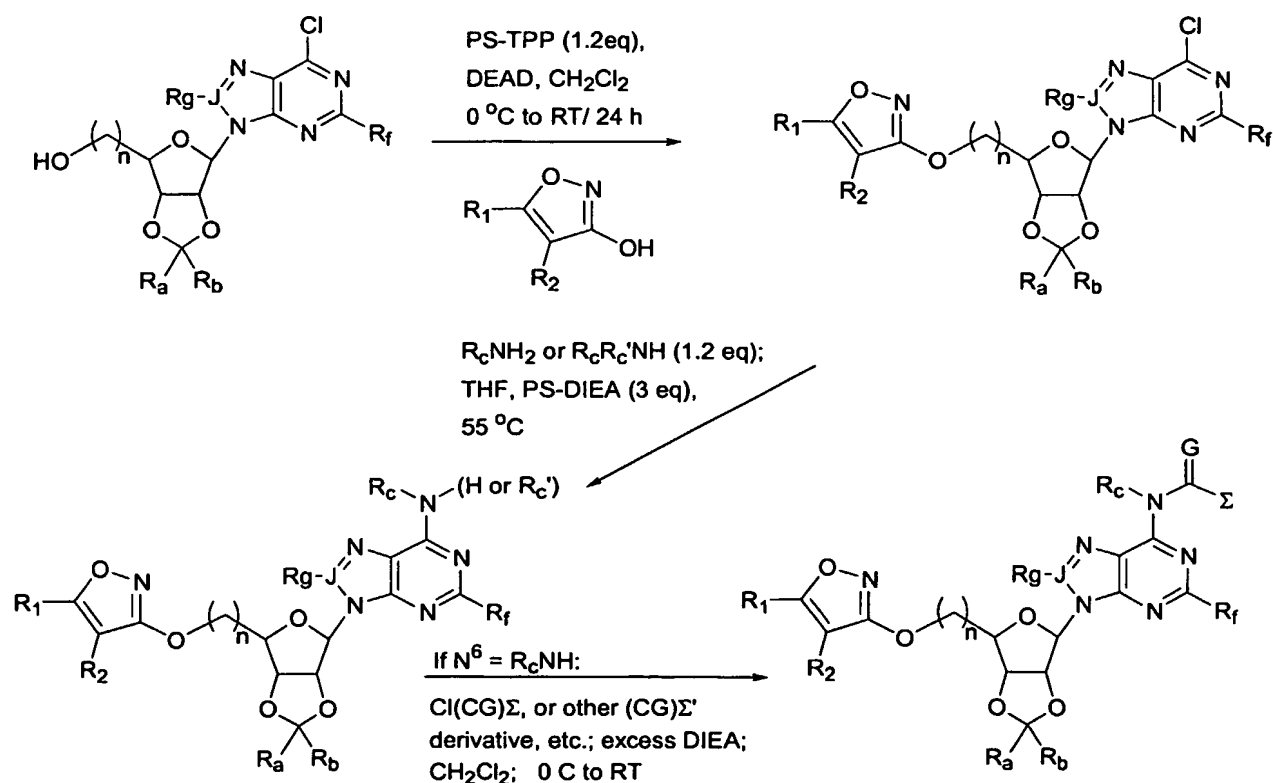
15 In addition to the introduction of amines on the C⁶ position using solid phase techniques as provided in Scheme 5, diversity can also be introduced at the 6-position of an adenosine, 8-azaadenosine, guanosine, etc., analog via the intermediacy of a 6-halogenated-purine derivative using solution phase methods. Scheme 6 exemplifies the preparation of 5'-isoxazole ethers, and shows the introduction of a nucleophile such as ammonia, an amine or a
20 hydroxylamine derivative at the 6-position of the purine/8-azapurine ring by displacement of a chloride leaving group (a 6-chloride is shown in Schemes 5 and 6, but the leaving group at C⁶ could also be another type, useful for such a transformation, e.g., a 6-bromide or 6-mesylate moiety) by such materials. Though Scheme 6 is exemplified using the generic primary amine R_cNH₂, a secondary amine such as disclosed for Scheme 5 could also be
25 employed, if desired, to give a N,N-disubstituted-N⁶-amine as a final product. N⁶-Substituted, and N⁶,N⁶-disubstituted amines are useful for the invention, as are N⁶-unsubstituted amines. Amines and amine-like compounds useful for displacement of a C⁶-halogen or another type of leaving group at C⁶-, as contemplated for these schemes comprise ammonia, methylamine, dimethylamine, and other N-alkyl- and N,N-dialkyl amines; N-
30 aralkylamines; N-cyclopropylamine and other N-cycloalkylamines; anilines; N-aryl-N-alkyl amines; ethers and other O-derivatives of hydroxylamine; aminopyridines and other heteroaromatic amines; heterocyclic compounds having a pendant -NR_cH group; heterocyclic amines, and alkyl amines which have one or more heteroatom units like O, NR, and/or S

substituted for carbon units in the alkyl chain. Such N⁶-products are useful in this invention. But, if desired, N⁶-products formed from said C⁶-halogen/leaving group displacements can be further transformed into amides, thioamides, ureas, thioureas, carbamates, guanidines, etc., by literature methods or by methods disclosed for such transformations in Scheme 5 and/or

5 Scheme 6 to give other derivatives useful in this invention. Following cleavage from the solid phase (for Scheme 5), the materials of Schemes 5 and/or Scheme 6 can be purified by methods typically used in the literature, such as by chromatography or, in certain cases, by crystallization. Preferred substituents at N⁶ are H, alkyl-groups, aryl-groups, amides, carbamates, and ureas.

10

Scheme 6. Solution phase synthesis of 5'-heteroaromatic ether-N⁶-derivatives by Mitsunobu reaction, 6-chloride displacement, and (if desired) N⁶-amine derivatization.

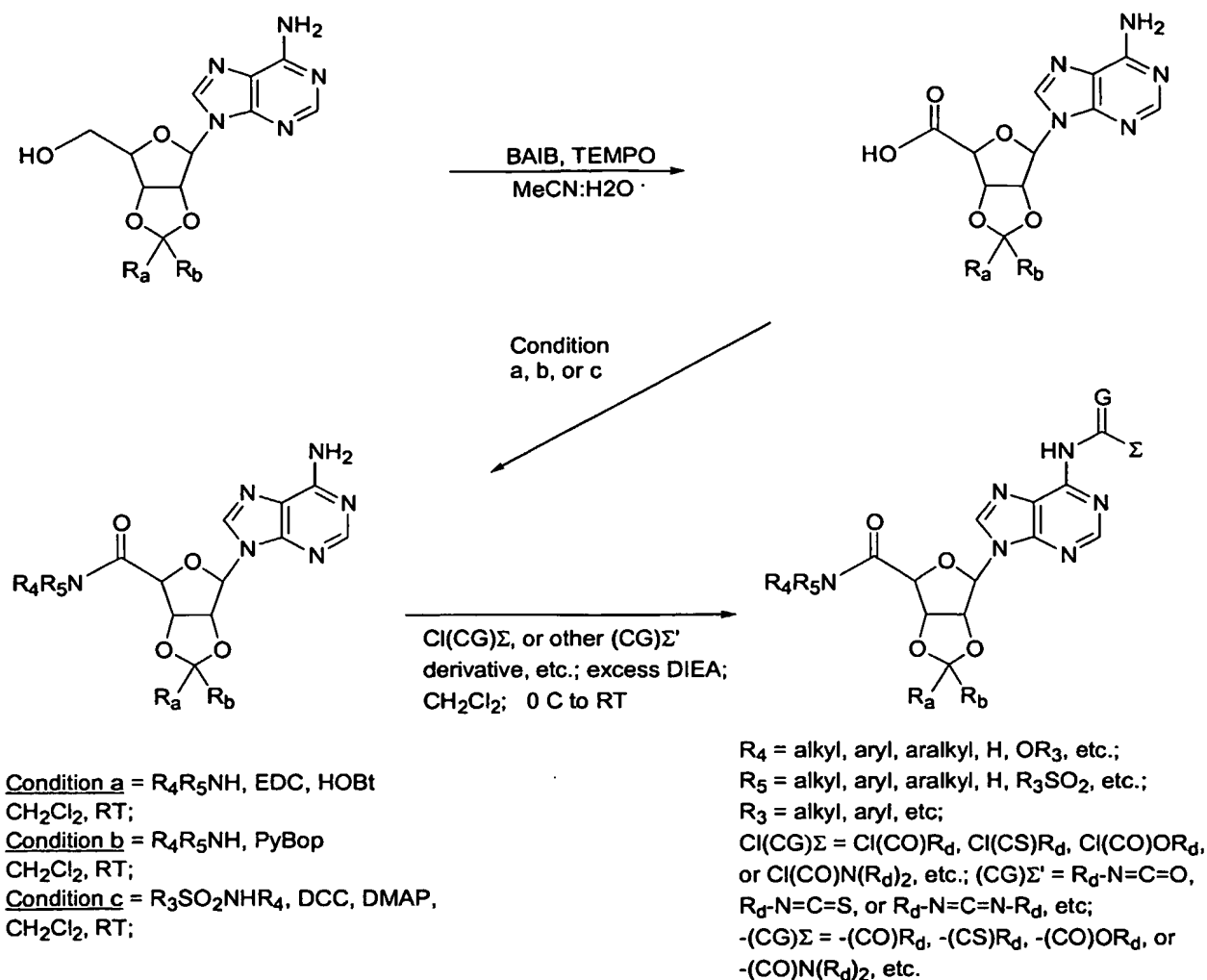


R₁, R₂ = CO₂R₃, alkyl, aryl, H, halogen, etc.
R₃ = alkyl, aryl etc; n = 1-6;
Cl(CG)Σ = Cl(CO)R_d, Cl(CS)R_d, Cl(CO)OR_d,
or Cl(CO)N(R_d)₂, etc.; (CG)Σ' = R_d-N=C=O,
R_d-N=C=S, or R_d-N=C=N-R_d, etc;
-(CG)Σ = -(CO)R_d, -(CS)R_d, -(CO)OR_d, or
-(CO)N(R_d)₂, etc.

15

When R_1 and/or R_2 of an isoxazole derivative as provided in Scheme 6 contains an ester group (e.g., $-C(CO)O-(alkyl)$, $-C(CO)O-(aryl)$, $-(CH_2)_mC(CO)O-(alkyl)$, $-O(CH_2)_mC(CO)O-(aryl)$, etc., where "m" defines the chain length of such a compound of Formula I), said ester can be used in the present invention, or it may be converted into an acid using a method which is compatible with an acetal or ketal moiety and a desired group at N^6 . For example, an ester provided in Scheme 6 can be hydrolyzed at room temperature (RT) in several hours using an excess of aqueous 2M lithium hydroxide solution dissolved in dioxane and/or methanol to give a carboxylate salt. Purification of said salt or a corresponding acid of said salt can be accomplished as previously disclosed. These acids and acid derivatives are also useful in this invention. Scheme 6 is exemplified using both a generic primary amine (which includes ammonia and hydroxylamine derivatives), and a generic secondary amine (including hydroxylamine derivatives) as choices for nucleophiles in the displacement of a leaving group (like chloride) at C^6 . As in Scheme 5, the synthesis of a C^6 -primary, secondary, or tertiary amine is possible using such a displacement reaction as shown in Scheme 6; such compounds are useful in this invention. Further modification of the N^6 -group of a compound of Scheme 6 with a $-(CG)\Sigma$ group or a $-(CG)\Sigma'$ group also yields a compound of the invention.

Scheme 7. Synthesis of 2', 3'-O-Isopropylideneadenosine-5'-carboxylic acid, related acetals and ketals, and their use as intermediates for the synthesis of compounds of this invention.

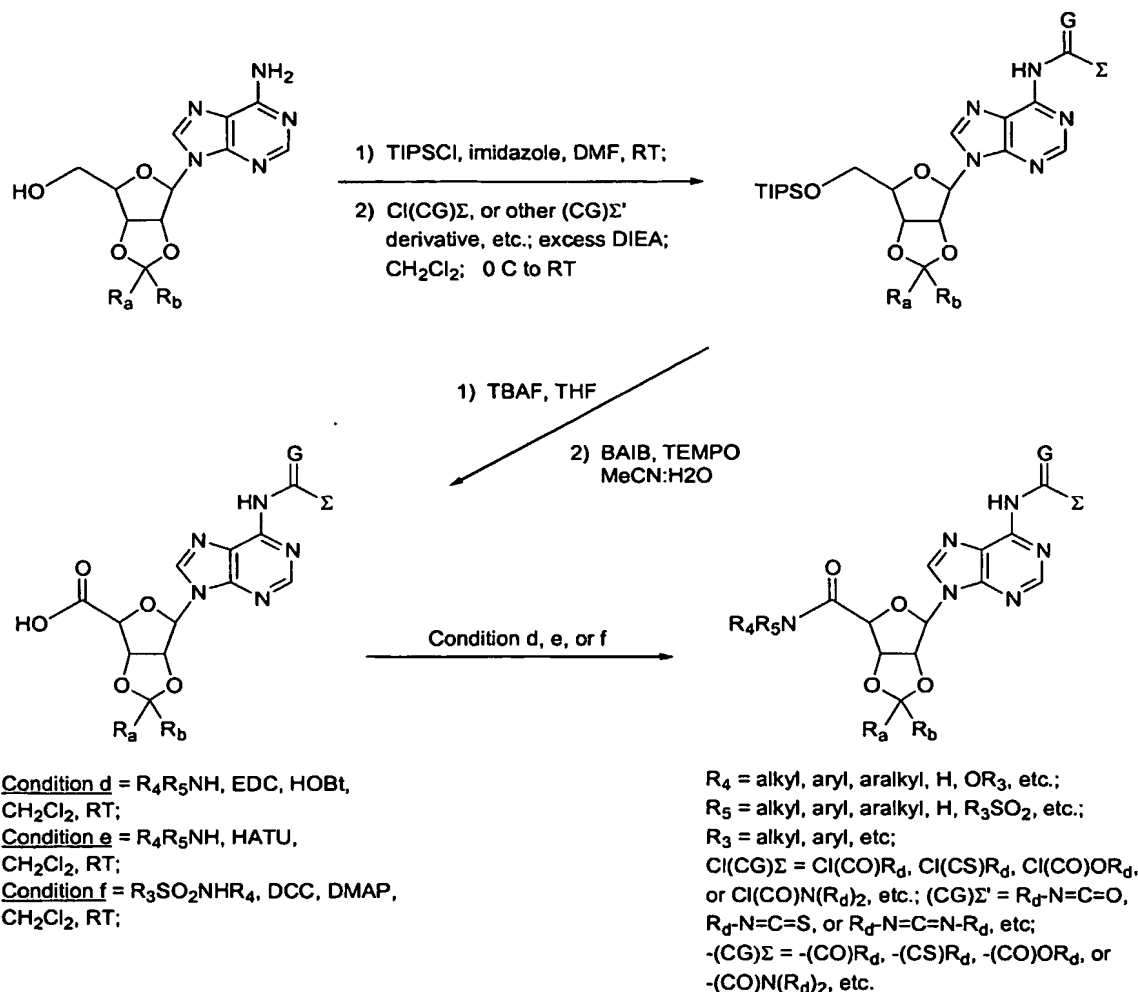


2', 3'-O-Isopropylideneadenosine-5'-carboxylic acid and related acetals and ketals are useful intermediates for the preparation of various amides and sulfonamide derivatives. An intermediate of this type can be used in both solution phase and solid phase synthetic schemes to prepare compounds with various substituents at the 5'-, 2', 3'-acetal/ketal-, and 6-positions- of adenosines or related purine derivatives as provided in Scheme 7. The methods shown in Scheme 7 are also useful for the preparation of substituted purine derivatives, and/or 8-azapurine derivatives.

Compounds containing the adenosine 5'-carboxylic acid unit shown in Scheme 7, or related N⁶-substituted-adenosine-5'-carboxylic acids and 8-azaadenosine-5'-carboxylic acids

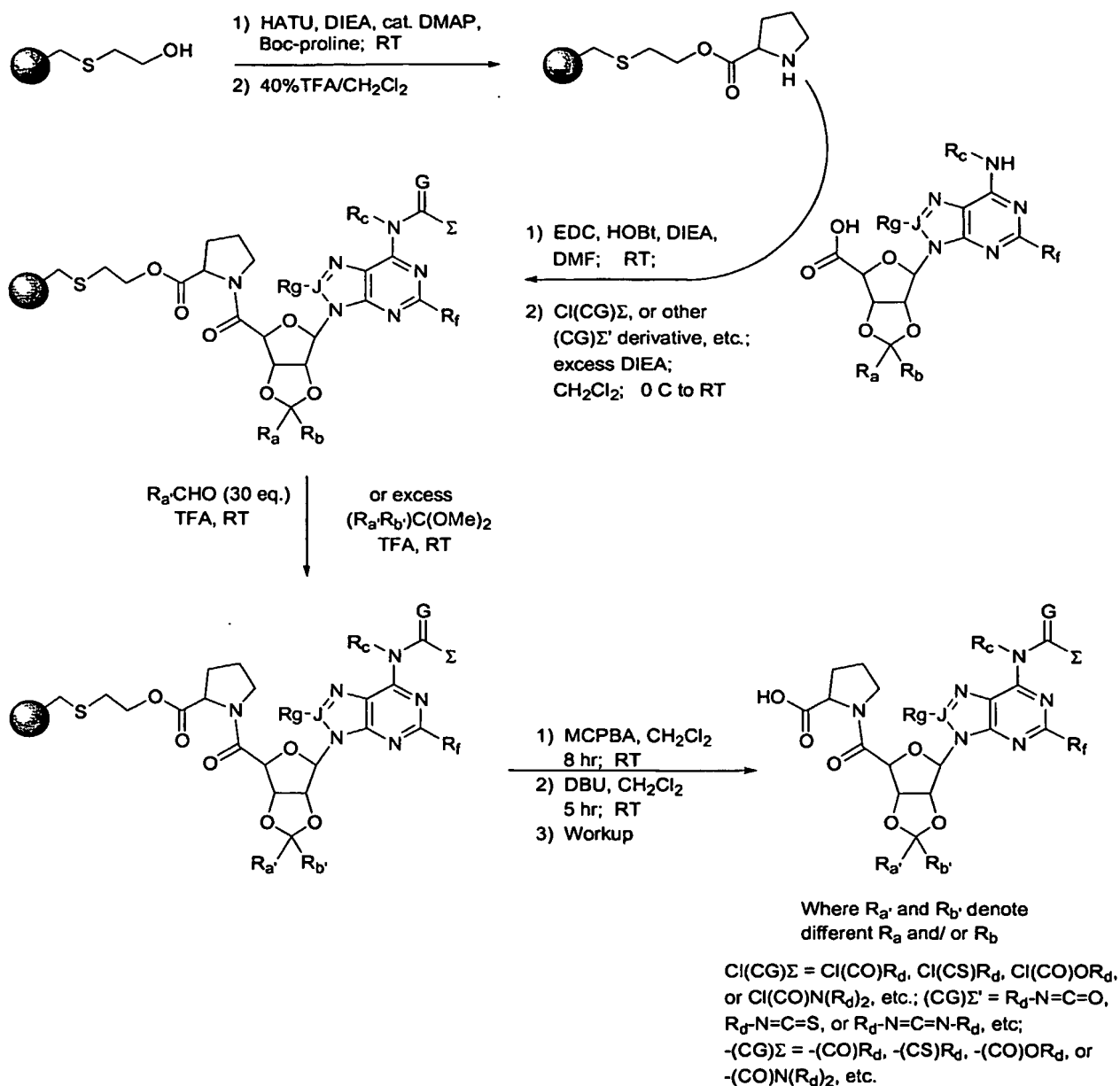
can also be used for the synthesis of esters or other materials useful in this invention using literature methods. In addition, the N⁶-amine of such an acid can be transformed into a urea, thiourea, amide, carbamate, guanidine, etc., to give a material that is also useful for this invention. Alternatively, a protecting group such as an amide (e.g., a benzamide, formamide, or trifluoroacetamide) or a carbamate (e.g., Boc, Cbz, Fmoc, etc.) can be installed on the N⁶ position, and then the acid moiety which is at, or linked to, the 5'-position of the furanose derivative can be coupled to a solid phase resin such as a 4-sulfamylbutyryl resin, hydroxymethyl resin, or a pegylated-hydroxy resin, etc., using techniques well-known in the literature to give a resin-bound material. The benefits of solid-phase chemistry can then be gained for modification of a resin-bound material by transketalization techniques similar to those shown in Schemes 4 and 5. If a protecting group is used on the N⁶-position of such an acid, it can be removed subsequent to a transketalization procedure to give a resin-bound compound which can be cleaved from the solid phase and purified to give a compound of the invention. Or, if desired, a N⁶-amine so formed on the solid phase can then be converted into an amide, carbamate, urea, thiourea or guanidine, etc., by methods disclosed herein, or by methods in the literature. Cleavage from the solid support using known methods then yields a compound of the invention which can be purified, if needed, by employing commonly-used techniques.

Scheme 8. N⁶-modification of adenosine derivatives, followed by oxidation to a 5'-carboxylic acid, and 5'-amide formation.



- 5 A variation of the method disclosed in Scheme 7, useful for the introduction of a group at N⁶ prior to oxidation of the 5'-position of an adenosine derivative or an 8-azaadenosine derivative, is shown in Scheme 8. Although Scheme 8 shows an adenine unit, it will be understood by chemistry practitioners that the methods of Scheme 8 are generally applicable to substituted members of the adenine family and also to the 8-azaadenines.
- 10 Preferred amines and amine derivatives for the 5'-amide-forming reactions shown in Schemes 7 and 8 are: trifluoromethanesulfonamide, methanesulfonamide, serine, glycine, proline, anthranilic acid and its regioisomers, and methyl anthranilate and its regioisomers.

Scheme 9. Amino acid amides of 5'-adenosinecarboxylic acid derivatives and 8-azaadenosine-5'-carboxylic acid derivatives using solid phase techniques.



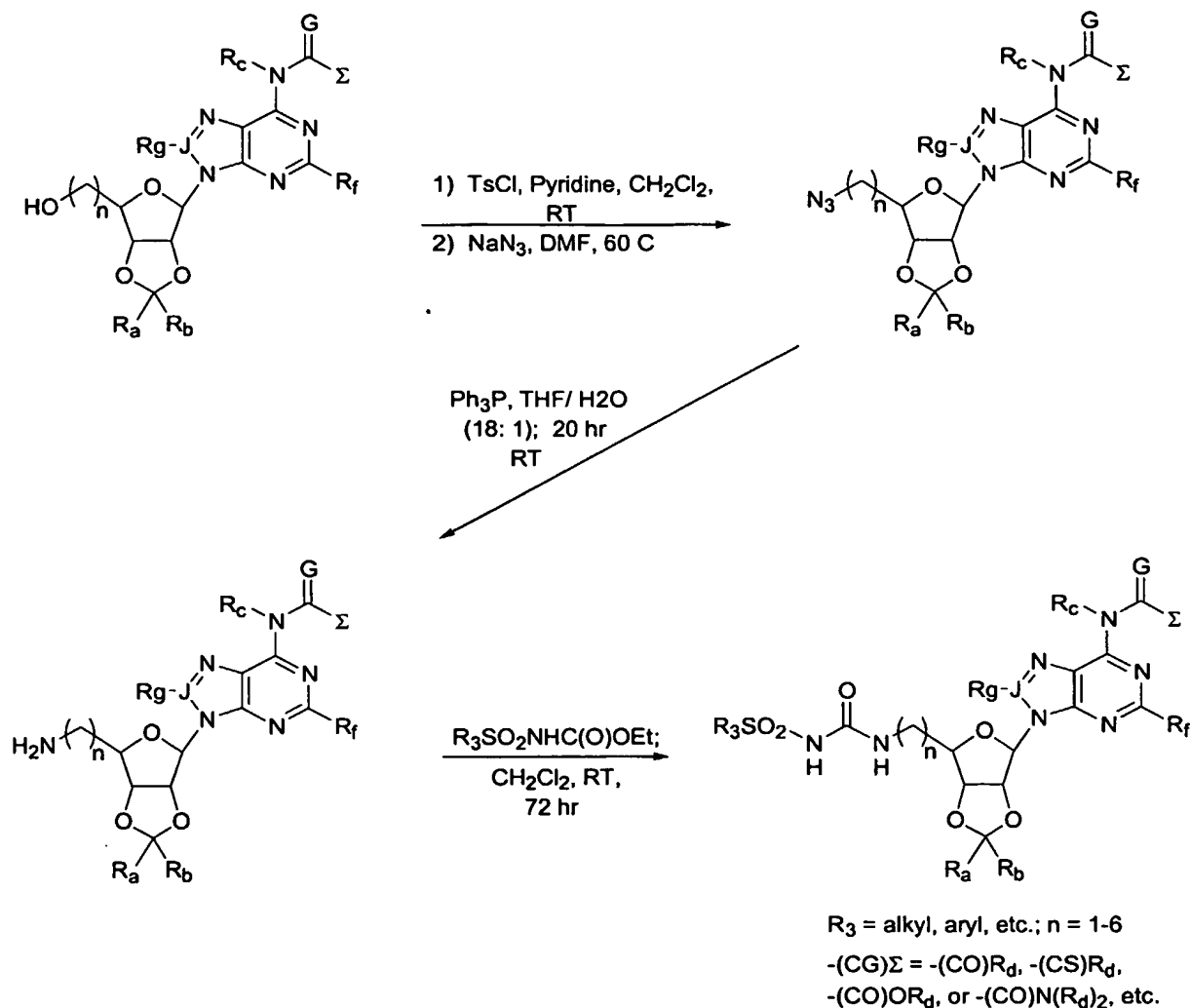
5

Amide derivatives of 5'-carboxylic acids (e.g., those shown in Scheme 7 and Scheme 8) or amide derivatives of acid moieties linked at the 5'-position of Formula I can also include amides derived from amino acids, peptides, amino alcohols and the like. A convenient way of attaching naturally-occurring, as well as synthetically-derived amino acids and peptides, or derivatives, to a 5'-carboxylic acid or related homologue is exemplified in Scheme 9 using the amino acid, proline.

10

An example of the method is shown in Scheme 9 utilizing a resin-linker combination like the polystyrene/HESM as a solid phase. Other solid phase/linkers known in the art can be used in this method. Attachment of a group, such as an amino acid, (e.g., proline, as shown) or a series of amino acids or a peptide, by well-known methods in the art of solid phase synthesis yields a resin-linked amine. Said amine can then be reacted with an adenosine 5'-carboxylic acid, or another type of derivative useful for making compounds of this invention, to yield a coupled product. The said coupled product, if it bears an amine at the 6-position, can then be treated with one of the various reagents described previously, or with a reagent known in the literature, to yield an amide, carbamate, urea, thiourea or guanidine, etc., at the adenosine 6-position. Alternatively, if a 5'-carboxylic acid derivative used in a coupling to a solid phase has a desired group already installed at the 6-position, then the latter said modification at N⁶ need not be performed. If desired, a coupled 5'-amide/N⁶-functionalized product can be converted into a variety of different acetals or ketals using solid phase methods such as described for Schemes 4 and 5. When a synthesis is complete, cleavage of a compound of the invention from a solid phase can be performed by a variety of methods known in the art; such cleavage conditions depending upon the type of linker used. Cleavage methodologies useful for cleaving peptide- or amino acid derivatives of 5'-linked-adenosine compounds comprise the linker oxidation/elimination procedures given in Schemes 5 and 9; treatment with a hydroxide source, such as lithium hydroxide, using conditions as for the ester hydrolysis described in Scheme 6; and hydrolysis using potassium trimethylsilanolate, as described in Scheme 4; as well as others known in the art (including aminolysis to form amides) which are useful for cleavage of other resin/linker/ester combinations. If desired, a compound useful in this invention can be obtained in purified form by cleaving it from a resin and purifying it as described previously.

Scheme 10. Synthesis of 5'-amines and 5'-amine derivatives of nucleosides and 8-azanucleosides.

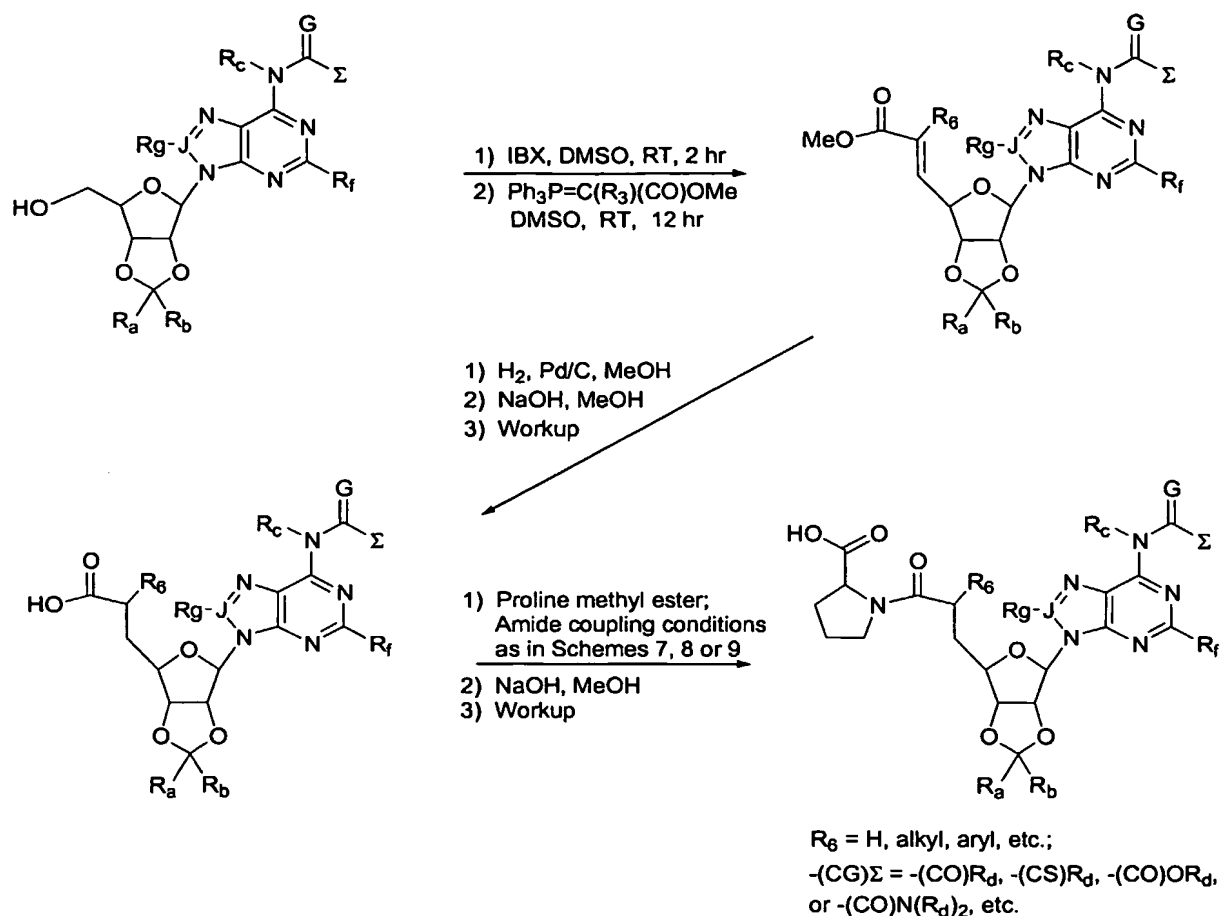


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In another embodiment, an amino group can be installed at the 5'-position of an adenosine or 8-azaadenosine analogue, or on the chain of a 5'-homologue of such a material. This amine can be utilized to form amide-, sulfonamide-, and other derivatives. Scheme 10 illustrates how a sulfonylurea can be synthesized at the 5'-position using a 5'-amine, or at related positions on homologous amine derivatives. In addition, an amine introduced at the 5'-position or on the 5'-chain of a homologue is also useful for the synthesis of amides, ureas, carbamates, sulfonamides and other amine derivatives using methods known in the art for such processes.

10

Scheme 11. Homologation of adenosine, including steps involving oxidation, Horner-Emmons Reaction, reduction, and coupling with amines.

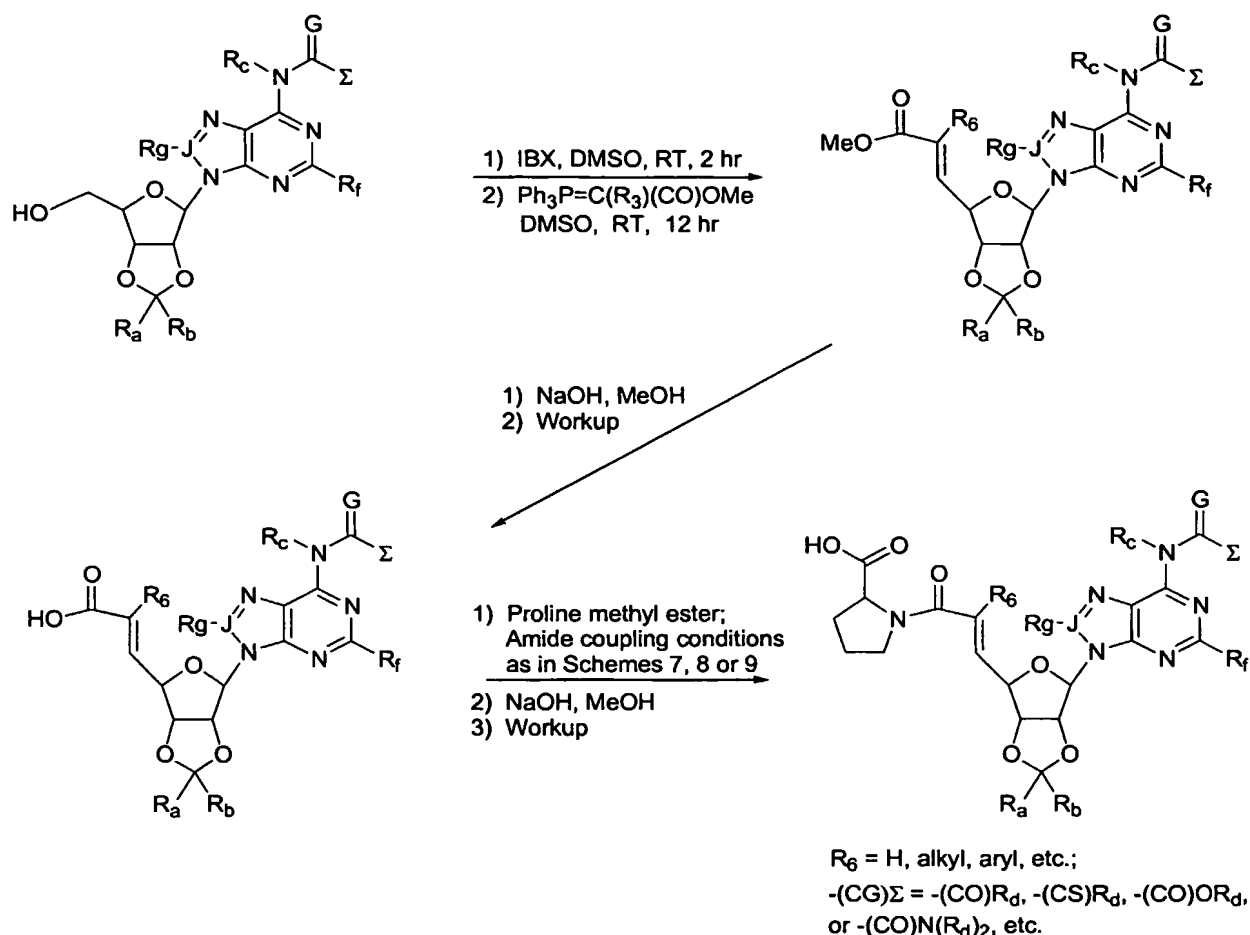


5

In still another embodiment, the 5'-position of the nucleoside derivative or 8-azanucleoside derivative is homologated with one or more carbon atoms. Scheme 11 illustrates the preparation of a class of homologated adenosine analogs which are useful for the invention. Or, if desired, such a homologue can subsequently be coupled with an amino acid to give other compounds useful for this invention. In Schemes 11 and 12, a proline is used to exemplify the amide coupling, but other amines or amino acid derivatives can be employed. In Scheme 12, the reduction step of Scheme 11 is omitted, which generates unsaturated homologues useful in this invention.

15

Scheme 12. Homologation of adenosine derivatives, including steps involving oxidation, Wittig or Horner-Emmons Reaction, and coupling with amines.



5

Those having skill in the art will recognize that the starting materials can be varied and additional steps employed to produce compounds encompassed by the present invention, as shown in the above schemes, as demonstrated by the examples which follow, and/or by appropriate modification of reactions known in the art. In some cases, protection of certain reactive functionalities may be necessary to achieve some of the above transformations. In general, the need for such protecting groups as well as the conditions necessary to attach and remove such groups will be apparent to those skilled in the art.

Methods of Administration

The active compounds disclosed herein are administered in a sufficient amount or amounts to reduce or alleviate the sensation of pain in a subject in need of such treatment by

either systemic administration, or direct administration to the painful area of a subject by any suitable means. In one embodiment, a compound of the invention is administered by administering a liquid or gel suspension of the active compound in the form of injection, drops, spray or gel. In a second embodiment, the active compound or compounds may be applied to the painful area via liposomes and/or using a cream formulation. In a third embodiment, the active compound or compound may be infused into the painful area via a pump-catheter system. In a fourth embodiment, one or more active compounds can be administered orally by any suitable means known in the art, but preferably said oral administration is in the form of a tablet, caplet, or capsule. Another embodiment of the present invention involves the active compound contained within a continuous or selective-release device, for example, membranes such as, but not limited to, those employed in the OCUSERT™ System (Alza Corp., Palo Alto, CA). As an additional embodiment, the active compound or compounds can be contained within, carried by, or attached to contact lenses that are placed on the eye, or adhesive bandages placed on the skin. Another embodiment of the present invention involves at least one active compound contained within or placed on a swab or sponge that can be applied to the painful area. Another embodiment of the present invention involves the active compound or compounds contained within a liquid spray that can be applied to the painful area. Such a spray also can be used for intranasal administration of one or more compounds of this invention such that one or more active materials contact a painful area of the nasal and/or sinus passages and reduce the sensation of pain in that area either by direct contact with the material or by systemic absorption. Other embodiments of the present invention involve an injection of at least one active compound directly into the painful area or application of at least one active compound directly or indirectly onto the surface of the painful area.

The quantity of each active compound of the invention included in a pharmaceutical composition of the present invention is an amount that is effective in reducing pain. The amount of each compound of the invention used is preferably an amount sufficient to achieve dissolved concentrations of each active compound in the painful area of the subject of from about 10^{-7} to about 10^{-1} moles/liter, and more preferably from about 10^{-6} to about 10^{-1} moles/liter, in order to antagonize the nociceptive effects of P2X receptors.

A topical solution comprising the active compound optimally contains a physiologically compatible vehicle, as those skilled in the art can select using conventional criteria. The vehicles are selected from the known ophthalmic, intranasal and cutaneous

vehicles which include, but are not limited to, saline solution, water-soluble polyethers such as polyethylene glycol, polyvinyls such as polyvinyl alcohol and povidone, cellulose derivatives such as methylcellulose and hydroxypropyl methylcellulose, petroleum derivatives such as mineral oil and white petrolatum, animal fats such as lanolin, polymers of acrylic acid
5 such as carboxypolymethylene gel, vegetable fats such as peanut oil and polysaccharides such as dextrans, and glycosaminoglycans such as sodium hyaluronate and salts such as sodium chloride and potassium chloride.

In addition to the topical method of administration described above, there are various methods of administering the active compounds of the present invention systemically. One
10 such means would involve an aerosol suspension of respirable particles comprising the active compound, which the subject inhales. The active compound would be absorbed into the bloodstream via the lungs, and subsequently affect a painful area in a pharmaceutically effective amount. The respirable particles may be liquid or solid, with a particle size sufficiently small to pass the mouth and larynx upon inhalation; in general, particles ranging
15 from about 1 to 10 microns, but more preferably 1-5 microns, in size are considered respirable.

Another means of systemically administering the active compounds to the eyes of the subject would involve administering a liquid/liquid suspension in the form of eye drops or eye wash or nasal drops of a liquid formulation, or a nasal spray of respirable particles that
20 the subject inhales. Liquid pharmaceutical compositions of the active compound for producing a nasal spray, or nasal or eye drops may be prepared by combining the active compound with a suitable vehicle, such as sterile pyrogen-free water or sterile saline by techniques known to those skilled in the art.

Other means of systemic administration of the active compound would involve oral
25 administration, in which pharmaceutical compositions containing active compounds are in the form of tablets, lozenges, aqueous or oily suspensions, dispersible powders or granules, emulsion, hard or soft capsules, or syrups or elixirs. Compositions intended for oral use can be prepared according to any method known to the art for the manufacture of pharmaceutical compositions and such compositions may contain one or more agents selected from the group
30 comprising sweetening agents, flavoring agents, coloring agents and preserving agents in order to provide pharmaceutically elegant and palatable preparations. Tablets contain the active ingredient in admixture with nontoxic pharmaceutically acceptable excipients that are suitable for the manufacture of tablets. These excipients include, for example, inert diluents,

such as calcium carbonate, sodium carbonate, lactose, calcium phosphate or sodium phosphate; granulating and disintegrating agents, for example, corn starch, or alginic acid; binding agents, for example, starch, gelatin or acacia; and lubricating agents, for example magnesium stearate, stearic acid or talc. The tablets can be uncoated or coated by known techniques to delay disintegration and absorption in the gastrointestinal tract and thereby provide a sustained action over a longer period. For example, a time delay material such as glyceryl monostearate or glyceryl distearate can be employed. Formulations for oral use can also be presented as hard gelatin capsules wherein the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin, or as soft gelatin capsules wherein the active ingredient is mixed with water or an oil medium, for example, peanut oil, liquid paraffin or olive oil.

Additional means of systemic administration of the active compound to the painful area of the subject would involve a suppository or pessary form of the active compound, such that a therapeutically effective amount of the compound reaches the painful area via contact or via systemic absorption and circulation.

Further means of systemic administration of the active compound would involve direct intra-operative instillation of a gel, cream, or liquid suspension form of a therapeutically effective amount of the active compound.

The method of the present invention can be used with other therapeutic and adjuvant agents commonly used to reduce pain, thus enhancing the effects of therapeutic agents and adjunctive agents. Other therapeutic agents used include opioids (morphine, fentanyl), sodium channel blockers (novocaine, lidocaine), NSAIDS (aspirin, ibuprofen), and COX-2 inhibitors (VIOXX®, CELEBREX®).

High doses are sometimes required for some therapeutic agents to achieve levels to effectuate the target response, but high doses are often associated with a greater frequency of dose-related adverse effects. Thus, combined use of one or more of the compounds of the present invention with therapeutic agents commonly used to treat pain allows the use of relatively lower doses of other agents, which results in a lower frequency of adverse side effects associated with long-term administration of such agents. Thus, another advantage of the compounds in this invention is to reduce adverse side effects of other drugs used to treat pain. Side effects known for other drugs, such as tolerance, dependence, constipation, respiratory depression, sedation, gastrointestinal side effects and increased bleeding time, can be reduced by administering a dose of one or more compounds of this invention to a subject

in need of pain reduction which thereby allows the concomitant reduction of a dose of another compound used to treat pain.

The invention is illustrated further by the following examples that are not to be construed as limiting the invention in scope to the specific procedures described in them.

5

EXAMPLES

Example 1. Preparation of 5'-aryl ether derivatives:

5-Amino-2-{2-benzyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4- 10 d][1,3]dioxol-4-ylmethoxy}-benzoic acid:

Adenosine (10 g, 37 mmol) was dissolved in N,N-dimethyl formamide (100 mL) and dimethoxypropane (25 mL) followed by addition of Amberlyst 15H⁺ resin. The mixture was stirred 3h at 55°C. The resin was removed by filtration and the solvents removed *in vacuo*, affording 2', 3'-di-O-isopropylidene adenosine (11 g, 95%).

15

This product (6 g, 20 mmol) was dissolved in N,N-dimethyl formamide (22 mL) and stirred with triisopropylsilyl chloride and imidazole 16 h at 23°C. The solution was partitioned between ether (200 mL) and brine (100 mL) and the ether phase washed with additional brine (2 x 50 mL). The ether was dried over magnesium sulfate and evaporated, affording 5'-O-triisopropylsilyl-2', 3'-di-O-isopropylidene adenosine.

20

This residue was dissolved in toluene (20 mL) and treated with phenylisocyanate (3.6 g, 30 mmol) for 16 h at 25°C. A solution of sodium bicarbonate (1 mL of 10 M) was added and the mixture evaporated to dryness. The residue was partitioned between ethyl acetate (100 mL) and water (25 mL). The organic phase was dried with magnesium sulfate and evaporated to dryness. The solid was dissolved in tetrahydrofuran (20 mL) and stirred with
25 tetrabutyl ammonium fluoride in tetrahydrofuran (20 mL of a 1 M solution) for 1 h in a dry ice/acetone bath. Removal of the solvent *in vacuo* followed by washing with hexane afforded the 5'-alcohol (5.3 g).

25

A portion of the above phenylurea product (0.41 g, 0.96 mmol) was suspended in 25 mL of 20% aqueous acetic acid and 5 mL of tetrahydrofuran /dioxane (1:1) and was stirred at
30 50°C for 24 h. The white suspension became a clear yellow solution. The mixture was concentrated and then lyophilized, to give 0.360 g (97% yield) of 1-[9-(3,4-dihydroxy-5-

hydroxymethyl-tetrahydro-furan-2-yl)-9H-purin-6-yl]-3-phenyl-urea as a yellow solid. MW calculated for $C_{17}H_{18}N_6O_5$ (MH^+) 387, found 387 by LCMS.

A small amount of 4A flame dried molecular sieves (cooled down by a flow of argon) was added to a vial containing a portion of the product immediately above (0.131 g, 0.34 mmol). The mixture was capped with a rubber septum and cooled down to 0 °C. To this mixture trifluoroacetic acid (2.5 mL) was added via syringe and the mixture stirred at this temperature for 15 min. Phenyl acetaldehyde dimethylacetal (0.230 mL, 4 eq.) was added dropwise and the mixture stirred at 0 °C for 2 h. One more equivalent of phenyl acetaldehyde dimethyl acetal was added and stirred an additional five hours. The volatiles were evaporated off and the residue was purified by flash chromatography (hexane: ethyl acetate, 8:2, 1% triethylamine) to give 0.095 g of product (60% yield) as a yellow solid. MW calculated for $C_{25}H_{24}N_6O_5$ (MH^+) 489, found 489 by LCMS..

A portion of this acetal product (0.068 g, 0.14 mmol) was dissolved in dry N,N-dimethyl formamide (2.5 mL) and potassium tert-butoxide (0.084 g, 5 eq) was added to give a yellow solution. To this mixture was added 2-fluoro-5-nitrobenzoic acid (0.046 g, 1.8 eq). After 2.5 h of stirring at room temperature the mixture was concentrated and purified by preparative HPLC to give the nucleoside analog as a white powder. MW calculated for $C_{32}H_{27}N_7O_9$ (MH^+) 654, found 654 by LCMS..

The nitro group of the product immediately above was reduced under a hydrogen atmosphere with a catalytic amount of 10% Pd/C in methanol during 6 h. Filtration through Celite followed by HPLC purification yielded 52 mg (62% yield) of the title compound as clear semisolid.

Example 2. Preparation of 5'-heteroaryl ether derivatives:

2-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-nicotinic acid:

To a stirred solution of 1-[9-(6-hydroxymethyl-2, 2-dimethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl)-9H-purin-6-yl]-3-phenyl-urea (43 mg, 0.1 mmol) in dry N, N'-dimethylformamide (1 mL) at 23 °C was added potassium *tert*-butoxide (45 mg, 0.4 mmol). After 30 min, 2-chloronicotinic acid (60 mg, 0.4 mmol) was added to the solution. The resulting mixture was stirred at 23 °C for 15 h and then quenched with water (1 mL), suspended in ethyl acetate (50 mL), washed with brine (3x), dried over anhydrous sodium

sulfate, filtered and concentrated *in vacuo* to give the crude product. Preparative reverse-phase HPLC was used to obtain the pure compound (15 mg, 27% yield) as a white solid. ¹H NMR (300 MHz, DMSO-d₆) δ 11.76 (s, 1H), 10.15 (s, 1H), 9.16 (s, 1H), 8.65 (s, 1H), 8.05 (dd, *J* = 8.5 Hz, 3.5 Hz, 1H), 7.94 (dd, *J* = 9.0 Hz, 3.0 Hz, 1H), 7.60 (d, *J* = 2.0 Hz, 1H), 7.57 (d, *J* = 2.0 Hz, 1H), 7.32 (t, *J* = 12.5 Hz, 2H), 7.05 (t, *J* = 12.5 Hz, 1H), 6.96 (dd, *J* = 12.5 Hz, 8.5 Hz, 1H), 6.25 (d, *J* = 6.0 Hz, 1H), 5.80 (t, *J* = 7.5 Hz, 1H), 5.10 (dd, *J* = 9.0 Hz, 2.5 Hz, 1H), 4.67 (m, 1H), 4.53 (dd, *J* = 20.0 Hz, 4.0 Hz, 1H), 4.36 (dd, *J* = 20.0 Hz, 4.0 Hz, 1H), 1.59 (s, 3H), 1.38 (s, 3H). MW calculated for C₂₆H₂₅N₇O₇ (MH⁺) 548, found 548 by LCMS..

10 Similarly, other 5'-substituted pyridines were prepared:

6-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-nicotinic acid: (9 mg, 8% yield) as a white solid. ¹H NMR (300 MHz, DMSO-d₆) δ 11.68 (s, 1H), 10.20 (s, 1H), 8.71 (s, 1H), 8.64 (d, *J* = 3.0 Hz, 2H), 8.12 (dd, *J* = 15.0 Hz, 3.5 Hz, 1H), 7.63 (d, *J* = 2.0 Hz, 1H), 7.61 (d, *J* = 1.5 Hz, 1H), 7.36 (t, *J* = 13.0 Hz, 2H), 7.09 (t, *J* = 12.5 Hz, 1H), 6.84 (dd, *J* = 13.5 Hz, 1.0 Hz, 1H), 6.33 (d, *J* = 3.5 Hz, 1H), 5.61 (dd, *J* = 10.0 Hz, 4.0 Hz, 1H), 5.20 (dd, *J* = 10.0 Hz, 4.5 Hz, 1H), 4.64 (m, 1H), 4.54 (m, 2H), 1.60 (s, 3H), 1.40 (s, 3H). MW calculated for C₂₆H₂₅N₇O₇ (MH⁺) 548, found 548 by LCMS..

20 **6-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-pyridine-2-carboxylic acid:** (5 mg, 4.6% yield) as a white solid. ¹H NMR (300 MHz, DMSO-d₆) δ 11.64 (s, 1H), 10.15 (s, 1H), 8.68 (s, 1H), 8.60 (s, 1H), 7.83 (dd, *J* = 13.5 Hz, 12.5 Hz, 1H), 7.62 (m, 3H), 7.34 (t, *J* = 12.5 Hz, 2H), 7.06 (t, *J* = 12.5 Hz, 1H), 6.96 (dd, *J* = 13.0 Hz, 1.0 Hz, 1H), 6.30 (d, *J* = 4.0 Hz, 1H), 5.59 (dd, *J* = 10.0 Hz, 4.5 Hz, 1H), 5.17 (dd, *J* = 10.0 Hz, 4.5 Hz, 1H), 4.64 (m, 1H), 4.50 (m, 2H), 1.58 (s, 3H), 1.38 (s, 3H). MW calculated for C₂₆H₂₅N₇O₇ (MH⁺) 548, found 548 by LCMS..

5-Chloro-6-{2,2-dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-nicotinic acid: (11 mg, 1.5% yield) as a white solid. MW calculated for C₂₆H₂₄ClN₇O₇ (MH⁺) 582, found 582 by LCMS..

6-Chloro-2-{2,2-dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-nicotinic acid (isomer A) & 2-Chloro-6-{2,2-dimethyl-6-[6-

(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-nicotinic acid (isomer B):

Two isomer were obtained from the reaction with 2, 6-dichloronicotinic acid. The major product was A (80 mg, 69% yield). LC-MS calculated for $C_{26}H_{24}ClN_7O_7$ (MH^+) 582, found 582. The minor product was B (20 mg, 17% yield). MW calculated for $C_{26}H_{24}ClN_7O_7$ (MH^+) 582, found 582 by LCMS..

2-Chloro-6-{2,2-dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-isonicotinic acid: (60 mg, 52% yield) as a white solid. 1H

NMR (300 MHz, DMSO- d_6) δ 11.63 (s, 1H), 10.18 (s, 1H), 8.68 (s, 1H), 8.61 (s, 1H), 7.6 (s, 1H), 7.59 (s, 1H), 7.36 (s, 1H), 7.33 (t, $J = 13.0$ Hz, 2H), 7.06 (t, $J = 13.5$ Hz, 1H), 6.29 (d, $J = 3.5$ Hz, 1H), 5.63 (dd, $J = 10.0$ Hz, 4.0 Hz, 1H), 5.17 (dd, $J = 10.0$ Hz, 4.5 Hz, 1H), 4.64 (m, 1H), 4.47 (m, 2H), 1.60 (s, 3H), 1.38 (s, 3H). MW calculated for $C_{26}H_{24}ClN_7O_7$ (MH^+) 582, found 582 by LCMS..

6-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-nicotinamide: (15 mg, 14% yield) as a white solid. 1H NMR (300 MHz,

DMSO- d_6) δ 11.70 (s, 1H), 10.10 (s, 1H), 8.69 (s, 1H), 8.61 (s, 1H), 8.60 (d, $J = 4.0$ Hz, 1H), 8.09 (dd, $J = 10.0$ Hz, 4.0 Hz, 1H), 7.95 (s, 1H), 7.62 (s, 1H), 7.59 (s, 1H), 7.39 (s, 1H), 7.34 (t, $J = 15.0$ Hz, 2H), 7.07 (t, $J = 12.5$ Hz, 1H), 6.80 (d, $J = 14.0$ Hz, 1.0 Hz, 1H), 6.31 (d, $J = 4.0$ Hz, 1H), 5.59 (dd, $J = 10.5$ Hz, 4.5 Hz, 1H), 5.17 (dd, $J = 10.0$ Hz, 4.5 Hz, 1H), 4.63 (m, 1H), 4.49 (m, 2H), 1.59 (s, 3H), 1.39 (s, 3H). MW calculated for $C_{26}H_{26}N_8O_6$ (MH^+) 547, found 547 by LCMS..

6-Chloro-2-{2,2-dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-5-fluoro-nicotinic acid: (35 mg, 19% yield) as a white solid. MW calculated for $C_{26}H_{23}ClFN_7O_7$ (MH^+) 600, found 600 by LCMS.

1-{9-[6-(3-Hydroxy-pyridin-2-yloxymethyl)-2,2-dimethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl]-9H-purin-6-yl}-3-phenyl-urea:

Dry argon was bubbled through a solution of 1-{9-[6-(3-Benzyloxy-pyridin-2-yloxymethyl)-2,2-dimethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl]-9H-purin-6-yl}-3-phenyl-urea (18 mg, 0.03 mmol, prepared as above) in methanol/ethyl acetate (1:1 v/v, 10 mL) at 23

°C for 10 min. Palladium on activated carbon (10%) was added and the suspension was degassed by argon for another 5 min. Hydrogen (H₂) was conducted to the solution via a balloon, and the reaction proceeded for 5h. The mixture was filtered and concentrated *in vacuo* to give the crude product. Preparative reverse-phase HPLC was used to obtain the pure compound (5 mg, 31% yield) as a white solid. MW calculated for C₂₅H₂₅N₇O₆ (MH⁺) 520, found 520 by LCMS.

2-{2-Benzyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-nicotinic acid:

2-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-nicotinic acid (180 mg, 0.33 mmol) was dissolved in a mixture of trifluoroacetic acid and water (TFA/H₂O, 4:1 v/v, 20 mL), and the suspension was stirred at 23 °C for 30 min. The solvents were removed under reduced pressure to give the intermediate diol product (120 mg, 72% yield) as a white solid. MW calculated for C₂₃H₂₁N₇O₇ (MH⁺) 508, found 508 by LCMS.

To a stirred solution of the diol immediately above (0.23 mmol) in dry trifluoroacetic acid (5 mL) at 23 °C was added phenyl acetaldehyde (130 mg, 1.1 mmol). The resulting mixture was stirred at 23 °C for 6h. After the removal of most trifluoroacetic acid by evaporation under reduced pressure, the reaction was quenched with saturated sodium bicarbonate solution (10 mL). The product was extracted using ethyl acetate (50 mL), washed with brine (3x), dried over anhydrous sodium sulfate, filtered and concentrated *in vacuo* to give the crude product. Preparative reverse-phase HPLC was used to obtain the pure acetal compound (7 mg, 5% yield) as a white solid. ¹H NMR (300 MHz, DMSO-d₆) δ 11.67 (s, 1H), 10.50 (s, 1H), 8.68 (s, 1H), 8.51 (s, 1H), 8.27 (dd, *J* = 8.0 Hz, 3.0 Hz, 1H), 8.10 (dd, *J* = 10.0 Hz, 4.0 Hz, 1H), 7.60 (dd, *J* = 14.0 Hz, 2.0 Hz, 1H), 7.30 (m, 7H), 7.08 (m, 3H), 6.21 (d, *J* = 4.5 Hz, 1H), 5.51 (dd, *J* = 10.5 Hz, 4.0 Hz, 1H), 5.32 (t, *J* = 8.0 Hz, 1H), 5.06 (dd, *J* = 10.0 Hz, 3.0 Hz, 1H), 4.70 (m, 1H), 4.62 (dd, *J* = 20.0 Hz, 5.0 Hz, 1H), 4.45 (dd, *J* = 20.0 Hz, 5.0 Hz, 1H), 3.10 (d, *J* = 8.5 Hz, 2H). MW calculated for C₃₁H₂₇N₇O₇ (MH⁺) 610, found 610 by LCMS.

Other 5'-substituted pyridine-ethers were transformed to various 2', 3'-acetals using methods similar to those given immediately above.

Example 3. Synthesis of 5'-isoxazole derivatives:**3-[6-(6-Chloro-purin-9-yl)-2,2-dimethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy]-isoxazole-5-carboxylic acid methyl ester:**

5 To a solution of [6-(6-Chloro-purin-9-yl)-2,2-dimethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl]-methanol (0.570 g, 1.74 mmol) in 16 mL of dry dichloromethane was added polymer-bound triphenylphosphine (PS-TPP; Argonaut Tech., 2.14 mmol/g, 0.91 g, 1.2 eq), followed by methyl-3-hydroxy-5-isoxazolecarboxylate (0.248 g, 1 eq). The mixture was sonicated for 15 minutes then stirred at room temperature for 1h under argon. The reaction
10 mixture was cooled to 0 °C and under argon flow diethylazodicarboxylate (0.33 g, 1.1 eq), dissolved in 1 ml of dichloromethane, was added dropwise via syringe. The mixture was protected from light and stirred at 0 °C for 30 min. then at room temperature for 20 h. The resin was washed liberally with dichloromethane and methanol. The organic solution obtained from the washes was concentrated to give, after flash chromatography purification,
15 0.740 g of the product as a white solid (95% yield). ¹H NMR (300 MHz, CDCl₃) δ 8.76 (s, 1H), 8.23 (s, 1H), 6.42 (s, 1H), 6.23 (d, *J* = 2.1 Hz, 1H), 5.43 (dd, *J* = 2.1 Hz, 1H), 5.14 (dd, *J* = 3.6 Hz, 1H), 4.7 (m, 1H), 4.61 (dd, *J* = 3.9 Hz, 1H), 4.49 (dd, *J* = 5.7 Hz, 1H), 3.93 (s, 3H), 1.65 (s, 3H), 1.42 (s, 3H). MW calculated for C₁₈H₁₈ClN₅O₇ (MH⁺) 452, found 452 by LCMS.

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3-{6-[6-(N-Benzyl-N-methyl-amino)-purin-9-yl]-2,2-dimethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-isoxazole-5-carboxylic acid:

To a solution of 3-[6-(6-Chloro-purin-9-yl)-2,2-dimethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy]-isoxazole-5-carboxylic acid methyl ester (0.106 g, 0.23 mmol) in
25 tetrahydrofuran (1.2 mL) was added polymer-bound diisopropylethylamine (PS-DIEA; Argonaut Tech., 3.83 mmol/g, 0.190 g, 3 eq), followed by addition of 0.040 mL of N-methyl-N-benzylamine (1.2 eq). The resulting mixture was stirred at room temperature overnight. The PS-DIEA resin was washed three times with dichloromethane and the solution obtained from the washes was concentrated to yield 3-{6-[6-(N-Benzyl-N-methyl-amino)-purin-9-yl]-
30 2,2-dimethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-isoxazole-5-carboxylic acid methyl ester as yellow oil. This material (0.23 mmol) was dissolved in 1,4-dioxane (1.2 mL) and 0.250 mL of an aqueous 2M lithium hydroxide solution added. The mixture was stirred at room temperature for 4 h. The crude product, isolated after acid workup, was used without

further purification in the following step. MW calculated for $C_{25}H_{26}N_6O_7$ (MH^+): 523, found 523 by LCMS.

3-{2-Benzyl-6-[6-(N-methyl-N-benzyl-amino)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-ylmethoxy}-isoxazole-5-carboxylic acid:

A solution of the above acetone (0.075 g, 0.140 mmol) dissolved in 1.5 mL of dry dichloromethane was cooled in an ice bath to 0 °C. To this clear mixture was added 1.8 mL of trifluoroacetic acid. The mixture was stirred at 0 °C for 5 h to afford the intermediate diol as a yellow semisolid after evaporative workup. This crude product (0.23 mmol) was dissolved at 0 °C in 1 mL of dry trifluoroacetic acid in the flask. To this mixture was added a small amount of 3A molecular sieves (previously dried by flame *in situ* and cooled by a stream of argon). The flask was capped with a rubber septum and cooled to 0 °C, then 0.1 mL of phenyl acetaldehyde dimethyl acetal was added and the mixture stirred for 18 h at 0 °C. At that point, 0.1 mL more of acetal was added and stirred an additional 6h. Purification by HPLC yielded 52 mg of the desired product as a clear semisolid (62% yield). MW calculated for $C_{30}H_{28}N_6O_7$ (MH^+): 585, found 585 by LCMS.

The following acetone-isoxazolecarboxylic acid analogs were prepared in a similar manner to the chloride displacement/hydrolysis sequence described above, using different amines:

Amine used; Product data:

- N-trans*-(cyclopropylphenyl)amine; Calculated mass for product (LCMS): $C_{26}H_{26}N_6O_7$ (MH^+): 535, found: 535.
- N*-Methyl-*N*-Phenethylamine; Calculated mass for product (LCMS): $C_{26}H_{28}N_6O_7$ (MH^+): 537, found: 537.
- Phenethylamine; Calculated mass for product (LCMS): $C_{25}H_{26}N_6O_7$ (MH^+): 523, found: 523.
- Benzylamine; Calculated mass for product (LCMS): $C_{24}H_{24}N_6O_7$ (MH^+): 509, found: 509.
- Isobutylamine; Calculated mass for product (LCMS): $C_{21}H_{26}N_6O_7$ (MH^+): 475, found: 475.
- Cyclopentylamine; Calculated mass for product (LCMS): $C_{22}H_{26}N_6O_7$ (MH^+): 487, found: 487.
- N*-(Methylthiophene-yl)amine; Calculated mass for product (LCMS): $C_{22}H_{22}N_6O_7S$ (MH^+): 515 found: 515.

5-Methylfurfurylamine; Calculated mass for product (LCMS): $C_{23}H_{24}N_6O_8$ (MH^+): 513, found: 513.

Propylamine; Calculated mass for product (LCMS): $C_{20}H_{24}N_6O_7$ (MH^+): 461, found: 461.

Aminopropylimidazole; Calculated mass for product (LCMS): $C_{23}H_{26}N_8O_7$ (MH^+): 527, found: 527.

3-Methylaminopyridine; Calculated mass for product (LCMS): $C_{23}H_{23}N_7O_7$ (MH^+): 510, found: 510.

Aminoethylpyrrolidine; Calculated mass for product (LCMS): $C_{23}H_{29}N_7O_7$ (MH^+): 516, found: 516.

10 Aniline; Calculated mass for product (LCMS): $C_{23}H_{22}N_6O_7$ (MH^+): 495, found: 495.

Example 4. Solution phase synthesis of 5'-carboxamide adenosine analogs

2', 3'-O-Isopropylideneadenosine-5'-carboxylic acid.

In a reaction vessel, bis-acetoxiodobenzene (BAIB, 1.15 g, 3.58 mmol), 2,2,6,6-tetramethyl-1-piperidinyloxy (TEMPO, 0.051 g, 0.325 mmol), and 2', 3'-isopropylideneadenosine (0.500 g, 1.63 mmol) were combined, and 3 mL of 1:1 acetonitrile:water mixture was added to the reaction vessel. The reaction mixture was stirred at ambient temperature under argon for 1 h then filtered. The white crystalline product was washed with acetonitrile:water mixture (1:1) and dried *in vacuo*, to yield 0.464 g of product, 89%. 1H NMR (300 MHz, CD_3SOCD_3) δ 12.77 (br s, 1 H), 8.22 (s, 1 H), 8.06 (s, 1 H), 7.27 (s, 2 H), 6.32 (s, 1 H), 5.52 (dd, $J_1 = 5.7$ Hz, $J_2 = 1.8$ Hz, 1 H), 5.45 (d, $J = 9.5$ Hz, 1 H), 4.67 (d, $J = 1.5$ Hz, 1 H), 1.52 (s, 3 H), 1.35 (s, 3 H).

2-({2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxole-4-carbonyl}-amino)-benzoic acid methyl ester:

To a vial containing 2', 3'-O-Isopropylideneadenosine-5'-carboxylic acid (0.241 g, 0.75 mmol) was added 3-amino-benzoic acid methyl ester (0.144 g, 0.75 mmol) in one portion at RT followed by heating at 50 °C overnight. The reaction mixture was diluted with 100 ml of ethyl acetate, washed with 1N hydrochloric acid, saturated sodium bicarbonate, brine, and dried over $MgSO_4$. Upon removal of solvent, the solid residue was purified with chromatography with 2-5 % methanol in dichloromethane to give 90 mg (26 %) of the carboxamide as white solid. MW calculated for $C_{21}H_{22}N_6O_6$ (MH^+) 455, found 455 by LCMS. This carboxamide product (90 mg, 0.20 mmol) was dissolved in DMF (2 mL) and

added to a flask containing phenyl isocyanate (35 mg, 0.30 mmol) in 2 ml of toluene at 50 °C. The reaction was stirred at 50 °C overnight. Additional phenyl isocyanate (35 mg, 0.20 mmol) was then added in several portions until nearly complete consumption of the starting material was noted by TLC analysis. The reaction mixture was then diluted with ethyl acetate (100 ml), washed with saturated sodium bicarbonate, brine, and dried over magnesium sulfate. The crude product was purified by a chromatography with 0-2 % methanol in dichloromethane to give 57 mg (50 %) of pure product and recovered starting material (10 mg). ¹H NMR (300 MHz, CDCl₃) δ 11.53 (s, 1H), 9.21 (s, 1H), 8.54 (s, 1H), 8.36 (s, 1H), 8.21 (s, 1H), 7.72 (m, 1H), 7.58 (m, 2H), 7.38 (m, 2H), 7.24 (m, 1H), 7.12 (m, 1H), 6.29 (d, *J* = 2.1 Hz, 1H), 5.66 (dd, *J* = 6.3, 1.5 Hz, 1H), 5.56 (dd, *J* = 6.3, 1.5 Hz, 1H), 4.88 (d, *J* = 1.8 Hz, 1H), 3.78 (s, 3H), 1.65 (s, 3H), 1.44 (s, 3H). MW calculated for C₂₈H₂₇N₇O₇ (MH⁺): 574, found 574 by LCMS.

3-({2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxole-4-carbonyl}-amino)-benzoic acid

To a vial containing 2', 3'-O-Isopropylideneadenosine-5'-carboxylic acid (250 mg, 0.778 mmol), triethylamine (1.57 mg, 1.56 mmol), and 3-amino-benzoic acid allyl ester (0.276 mg, 1.56 mmol) in 0.5 ml of N,N-dimethyl formamide at 0 °C was added PyBOP (0.443 mg, 0.856 mmol) in one portion. Reaction was continued at 0 °C for 4 h and at ambient for 4 h. Additional PyBOP (50 mg) was added and the reaction continued overnight at ambient temperature. The reaction mixture was diluted with ethyl acetate (100 ml), washed with saturated sodium bicarbonate, brine, and dried over magnesium sulfate. The crude was purified by a chromatography with 0-2 % methanol in dichloromethane to give the desired amide product. MW calculated for C₂₃H₂₄N₆O₆ (MH⁺) 481, found 481 by LCMS.

Phenyl isocyanate was coupled with the amide product using a method similar to those described above, affording the intermediate phenylurea compound. MW calculated for C₃₀H₂₉N₇O₇ (MH⁺) 600, found 600 by LCMS.

This phenylurea compound (36 mg, 0.057 mmol) and morpholine (0.015 mg, 0.17 mmol) were dissolved in tetrahydrofuran (5 ml) followed by addition of tetrakis(triphenylphosphine)palladium (5 mg, 0.004 mmol). Reaction was completed in 4 h at RT. After removal of the solvent, the crude product mixture was purified by preparative HPLC to give the desired product. ¹H NMR (300 MHz, CDCl₃) δ 11.52 (s, 1H), 9.99 (s, 1H),

9.60 (s, 1H), 8.61 (s, 1H), 8.42 (s, 1H), 7.94 (s, 1H), 7.72 (m, 4H), 7.34 (m, 2H), 7.23 (m, 1H), 7.06 (m, 1H), 6.56 (s, 1H), 5.56 (m, 2H), 4.88 (s, 1H), 1.58 (s, 3H), 1.39 (s, 3H).

Example 5. Solution phase synthesis of a sulfonamide:

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N-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxole-4-carbonyl}-methanesulfonamide:

To a vial containing 2', 3'-O-Isopropylideneadenosine-5'-carboxylic acid (20 mg, 0.045 mmol), dimethylaminopyridine (5 mg, 0.045 mmol), and methanesulfonamide (0.009 mg, 0.091 mmol) in dichloromethane (0.5 ml) was added dicyclohexylcarbodiimide (10 mg, 0.05 mmol). The mixture was stirred for 2 days at RT. Additional dicyclohexylcarbodiimide (10 mg, 0.05 mmol) and dimethylaminopyridine (5 mg, 0.045 mmol) were added and the reaction was continued at RT overnight. To the reaction mixture was added ethyl acetate (75 ml), which was washed with 1 N hydrochloric acid, water, saturated sodium bicarbonate, brine, and dried over magnesium sulfate. Upon removal of solvent, the residue was purified by prep. HPLC to yield 13 mg of desired product (55%). MW calculated for C₂₁H₂₃N₇O₇S (MH⁺) 518, found 518 by LCMS.

Example 6. Solid phase synthesis of ureas and acetals from polymer bound 5'-proline-amides of adenosine:

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Commercially available hydroxymethylsulfanylmethyl (HESM) polystyrene resin (1.4 mmol/g, 200 mesh, NovaBiochem; 2.82 g, 3.95 mmol) was swelled for 15 minutes in 50 mL. In a separate reaction vessel, Boc-Pro-OH (3.40 g, 15.8 mmol), HATU (5.7 g, 15.0 mmol), dimethylaminopyridine (0.24 g, 1.98 mmol), and diisopropylethylamine (3.5 mL, 19.8 mmol) were dissolved in 40 mL of N,N-dimethyl formamide and stirred for 15 minutes. N,N-dimethyl formamide was drained from the HESM resin and the solution of activated proline derivative was added to the resin. The resin was agitated at RT for 17 h. The solvent was then drained and the resin washed with N,N-dimethyl formamide (3 x 30 mL), dichloromethane (3 x 30 mL), methanol (3 x 30 mL), dichloromethane (2 x 30 mL), methanol (3 x 30 mL) and dried *in vacuo* overnight. Mass of resin: 3.42 g, 93% loading.

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Removal of BOC protecting group:

Resin obtained in the previous step was agitated with a 40% trifluoroacetic acid/dichloromethane solution (75 mL) for 15 minutes. The solvent was drained, and a fresh solution of 40% trifluoroacetic acid in dichloromethane was added, and resin was agitated for another 15 minutes. After this the resin was washed with dichloromethane (5 x 40 mL), 20% diisopropylethylamine/dichloromethane (2 x 30 mL), dichloromethane (3 x 30 mL), and methanol (5 x 40 mL). The resin was dried under vacuum. A chloranil test indicated the presence of a free amino group, and this proline-bound resin was carried over to the next step.

The proline resin product from the previous step was swelled in 50 mL N,N-dimethyl formamide for 30 minutes, after that the N,N-dimethyl formamide was drained. In a separate reaction vessel, 2', 3'-O-Isopropylideneadenosine-5'-carboxylic acid (1.40 g, 4.35 mmol), dichloroethane (0.91 g, 4.74 mmol), HOBT·H₂O (0.73 g, 4.74 mmol), and diisopropylethylamine (3.5 mL, 19.8 mmol) were dissolved in 55 mL of N,N-dimethyl formamide. The solution was stirred for 15 minutes, and then added to the proline resin. The resin was agitated at RT for 17 h. The solvent was drained, the resin was washed with N,N-dimethyl formamide (3x30 mL), dichloromethane (3x30 mL), methanol (3x30 mL), dichloromethane (2x30 mL), methanol (3x30 mL), and dried *in vacuo* for 48 h. A chloranil test performed on a few sample beads indicated that coupling had occurred. A small amount of resin was cleaved using the following procedure to verify attachment of the carboxylic acid, and analysed by LCMS. Calculated mass for C₁₈H₂₂N₆O₆ (MH⁺): 419, found 419 by LCMS. Mass of resin: 3.66 g, 0.55 mmol/g, 82% in three steps.

General cleavage procedure for analysis of HESM resin:

A small amount of resin is suspended in a solution of 5 - 6 equivalents of m-chloroperbenzoic acid in dichloromethane and agitated for 7 - 8 hours at RT. The solution is then drained, and the resin is washed 5 - 6 times with fresh dichloromethane. Then resin is suspended in a solution of 4 - 5 equivalents of DBU in dichloromethane, and agitated at RT for 4 - 5 hours. The resin is then filtered and the solution is analyzed by LC/MS, HPLC or another method. The compounds are recovered from solution by rotary evaporation.

The adenosine-proline amide-derivatized resin from the previous resin-synthesis step (0.5 g, 0.275 mmol) was suspended in anhydrous N,N-dimethyl formamide (10 mL). Ethyl isocyanate was added (0.43 mL, 5.5 mmol), and the reaction mixture was heated in a capped vial at 55 °C for 16 h. The resin was then drained and washed with N,N-dimethyl formamide

(3 x 10 mL), dichloromethane (3 x 10 mL), methanol (3 x 10 mL), dichloromethane (2 x 10 mL), methanol (3 x 10 mL), and the procedure was repeated once again then the resin dried *in vacuo* for 24 h. A negative chloranil test indicated complete reaction. A small amount of this resin (bound to the 5'-adenosine-(2', 3'-acetone)-proline amide-derivatized as the 6-ethylurea) was cleaved by the procedure described for cleavage of HESM resin, and the isolated product was analyzed by LCMS. Mass calculated for C₂₁H₂₇N₇O₇ (MH⁺): 490, found 490 by LCMS. Loading 0.52 mmol/g, mass 0.52 g, 98%.

The procedure described above was also used to prepare N⁶-ureas with R_d = -C₆H₁₁, -Ph, -CH₂Ph, -CH₂CH₂Ph, -cyclopentyl, and *trans*-2-phenyl-cycloprop-1-yl groups using the appropriate isocyanates.

1-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-phenyl-tetrahydro-furo[3,4-d][1,3]dioxole-4-carbonyl}-pyrrolidine-2-carboxylic acid:

The resin described above (0.05 g, 0.026 mmol) was suspended in trifluoroacetic acid, then benzaldehyde (0.095 g, 0.9 mmol) was added all at once. The resin was agitated in a tightly closed vial for 24 h. The resin was drained, washed with dichloromethane (5 x 3 mL), 20% diisopropylethylamine/ dichloromethane (2 x 3 mL), dichloromethane (3 x 3 mL), and methanol (5 x 3 mL), then dried *in vacuo* for 3 h. The resin was cleaved using the cleavage procedure described above, the crude product was collected, analyzed by LCMS and purified by preparative HPLC. Calculated MW for C₂₅H₂₇N₇O₇ (MH⁺): 538, found: 538 by LCMS.

The following analogs were prepared in a similar manner to that described above, using appropriate combinations of isocyanates and aldehydes. Compounds analyzed by LCMS.

R_d = ethyl, R_a = benzyl Calculated MW for C₂₆H₂₉N₇O₇: 552.55 (MH⁺), found: 552.4.

R_d = ethyl, R_a = 4-biphenyl Calculated MW for C₃₁H₃₁N₇O₇: 614.62 (MH⁺), found: 614.3

R_d = ethyl, R_a = 3-biphenyl Calculated MW for C₃₁H₃₁N₇O₇: 614.62 (MH⁺), found: 614.3

R_d = ethyl, R_a = 2-naphthyl Calculated MW for C₂₉H₂₉N₇O₇: 588.58 (MH⁺), found: 588.1.

R_d = n-hexyl, R_a = phenyl Calculated MW for C₂₉H₃₅N₇O₇: 594.63 (MH⁺), found: 594.3.

R_d = n-hexyl, R_a = benzyl Calculated MW for C₃₀H₃₇N₇O₇: 608.65 (MH⁺), found: 608.2.

R_d = n-hexyl, R_a = 4-biphenyl Calculated MW for C₃₅H₃₉N₇O₇: 670.73 (MH⁺), found:

670.3.

R_d = n-hexyl, R_a = 3-biphenyl Calculated MW for C35H39N7O7: 670.73 (MH+), found: 670.3.

R_d = n-hexyl, R_a = 2-naphthyl Calculated MW for C33H37N7O7: 644.69 (MH+), found: 644.3.

R_d = cyclopentyl, R_a = benzyl Calculated MW for C29H33N7O7: 592.62 (MH+), found: 592.3.

R_d = cyclopentyl, R_a = phenyl Calculated MW for C28H31N7O7: 578.59 (MH+), found: 578.3.

R_d = cyclopentyl, R_a = 4-biphenyl Calculated MW for C34H35N7O7: 654.68 (MH+), found: 654.3.

R_d = cyclopentyl, R_a = 3-biphenyl Calculated MW for C34H35N7O7: 654.68 (MH+), found: 654.3.

R_d = cyclopentyl, R_a = 2-naphthyl Calculated MW for C32H33N7O7: 628.65 (MH+), found: 628.4.

R_d = benzyl, R_a = benzyl Calculated MW for C31H31N7O7: 614.62 (MH+), found: 614.3.

R_d = benzyl, R_a = phenyl Calculated MW for C30H29N7O7: 600.59 (MH+), found: 600.3.

R_d = benzyl, R_a = 2-naphthyl Calculated MW for C34H31N7O7: 650.65 (MH+), found: 650.3.

R_d = benzyl, R_a = 4-biphenyl Calculated MW for C36H33N7O7: 676.69 (MH+), found: 676.3.

R_d = benzyl, R_a = 3-biphenyl Calculated MW for C36H33N7O7: 676.69 (MH+), found: 676.3.

R_d = ethylphenyl, R_a = benzyl Calculated MW for C32H33N7O7: 628.65 (MH+), found: 628.4.

R_d = ethylphenyl, R_a = phenyl Calculated MW for C31H31N7O7: 614.62 (MH+), found: 614.5.

R_d = ethylphenyl, R_a = 4-biphenyl Calculated MW for C37H35N7O7: 690.72 (MH+), found: 690.4.

R_d = ethylphenyl, R_a = 3-biphenyl Calculated MW for C37H35N7O7: 690.72 (MH+), found: 690.5.

R_d = ethylphenyl, R_a = 2-naphthyl Calculated MW for C35H33N7O7: 664.68 (MH+), found: 664.4.

R_d = cyclopropyl-*trans*-2-phenyl, R_a = benzyl Calculated MW for C33H33N7O7: 640.66 (MH+), found: 640.3.

R_d = cyclopropyl-*trans*-2-phenyl, R_a = 3-biphenyl Calculated MW for C38H35N7O7: 702.73

(MH+), found: 702.6.

R_d = cyclopropyl-*trans*-2-phenyl, R_a = 2-naphthyl Calculated MW for C₃₆H₃₃N₇O₇: 676.69 (MH+), found: 676.6.

R_d = phenyl, R_a = benzyl Calculated MW for C₃₀H₂₉N₇O₇: 600.59 (MH+), found: 600.3.

5 R_d = phenyl, R_a = phenyl Calculated MW for C₂₉H₂₇N₇O₇: 586.57 (MH+), found: 586.2.

R_d = phenyl, R_a = 2-naphthyl Calculated MW for C₃₃H₂₉N₇O₇: 636.63 (MH+), found: 636.5.

R_d = phenyl, R_a = 4-biphenyl Calculated MW for C₃₅H₃₁N₇O₇: 662.66 (MH+), found: 662.5.

10 R_d = phenyl, R_a = 3-biphenyl Calculated MW for C₃₅H₃₁N₇O₇: 662.66 (MH+), found: 662.5.

R_d = phenyl, R_a = 3-thianaphthene Calculated MW for C₃₁H₂₇N₇O₇S: 642.65 (MH+), found: 642.0.

15 R_d = ethylphenyl, R_a = 3-thianaphthene Calculated MW for C₃₃H₃₁N₇O₇S: 670.71 (MH+), found: 670.0.

R_d = ethylphenyl, R_a = 3-thianaphthene Calculated MW for C₃₂H₂₉N₇O₇S: 656.68 (MH+), found: 656.1.

R_d = n-hexyl, R_a = 3-thianaphthene Calculated MW for C₃₁H₃₅N₇O₇S: 650.72 (MH+), found: 650.2.

20 R_d = n-hexyl, R_a = CHCHPh Calculated MW for C₃₁H₃₇N₇O₇: 620.67 (MH+), found: 620.4.

R_d = n-hexyl, R_a = CCPh Calculated MW for C₃₁H₃₅N₇O₇: 618.65 (MH+), found: 618.0.

R_d = cyclopropyl-*trans*-2-phenyl, R_a = CCPh Calculated MW for C₃₄H₃₁N₇O₇: 650.65 (MH+), found: 650.1.

25 R_d = ethylphenyl, R_a = CCPh Calculated MW for C₃₃H₃₁N₇O₇: 638.64 (MH+), found: 638.1.

R_d = benzyl, R_a = CCPh Calculated MW for C₃₂H₂₉N₇O₇: 624.62 (MH+), found: 624.1.

R_d = ethyl, R_a = CCPh Calculated MW for C₂₇H₂₇N₇O₇: 562.55 (MH+), found: 562.0.

R_d = cyclopropyl-*trans*-2-phenyl, R_a = 4-biphenyl Calculated MW for C₃₈H₃₅N₇O₇: 702.73 (MH+), found: 702.1.

30 R_d = cyclopropyl-*trans*-2-phenyl, R_a = phenyl Calculated MW for C₃₂H₃₁N₇O₇: 626.63 (MH+), found: 626.0.

R_d = cyclopropyl-*trans*-2-phenyl, R_a = CHCHPh Calculated MW for C₃₄H₃₃N₇O₇: 652.67 (MH+), found: 652.1.

R_d = ethylphenyl, R_a = CHCHPh Calculated MW for $C_{33}H_{33}N_7O_7$: 640.66 (MH⁺), found: 640.3.

Example 7. (See Scheme 8) 1-[9-(2-Benzyl-6-ureidomethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl)-9H-purin-6-yl]-3-phenyl-urea methylsulfonamide:

In a reaction vessel were combined 2',3'-O-benzylidene-N6-(phenylurea)adenosine (0.200 g, 0.409 mmol), p-toluene sulfonyl chloride (0.148 g, 0.778 mmol), and 4.0 mL of pyridine. The mixture was stirred for 4 h at RT. The solvent was removed *in vacuo* and the tosylate (0.260 g, 99%) was recovered as a yellow solid. MW calculated for $C_{32}H_{30}N_6O_7S$: 643.19 (MH⁺), found 642.9.

This tosylate product (0.260 g, 0.409 mmol) was dissolved in 3 mL of anhydrous N,N-dimethyl formamide, sodium azide (0.266 g, 4.09 mmol) was added, and the mixture was heated at 80 °C in a closed vial for 7 h while being stirred. The mixture was diluted with 50 mL of dichloromethane and extracted with 5% sodium bicarbonate solution and brine. The organic layer was separated, dried over anhydrous sodium sulfate and solvent was removed *in vacuo*. The azide derivative was recovered (0.183 g, 87%) as a white solid. MW calculated for $C_{25}H_{23}N_9O_4$: 514.19 (MH⁺), found 514.1.

This residue containing the azide (0.180 g, 0.351 mmol) was dissolved in 6 mL of tetrahydrofuran/water mixture (18:1), polystyrene-bound triphenylphosphine was added (2.19 mmol/g, 0.800 g, 1.75 mmol), and the reaction mixture was stirred at RT under argon for 24 h. The reaction mixture was then filtered, the solvent was removed *in vacuo* and the crude product was chromatographed on a silica gel column (2 cm x 15 cm). The column was eluted with dichloromethane/methanol/triethylamine (88:10:2) to give 0.072 g (42%) of the amine as a white solid. MW calculated for $C_{25}H_{25}N_7O_4$: 488.20 (MH⁺), found 488.0.

A portion of the amine product above (0.022 g, 0.046 mmol) was dissolved in 2 mL of dichloromethane (anhydrous) and methylsulfonylethylcarbamate (0.008 g, 0.046 mmol) was added. The reaction was stirred at RT under argon for 72 h. Solvent was then removed *in vacuo*, and the crude product was purified by preparative HPLC (acetonitrile/0.1% trifluoroacetic acid //water/0.1% trifluoroacetic acid buffer). 0.016 g (59%) of product was recovered as a white solid. MW calculated for $C_{27}H_{28}N_8O_7S$: 609.18 (MH⁺), found 609.4. ¹H NMR (300 MHz, CD₃SOCD₃) δ 11.69 (br s, 1 H), 10.22 (br s, 1 H), 8.69 (s, 1 H), 8.65 (s, 1 H), 7.60 (d, *J* = 7.8 Hz, 2 H), 7.36 – 7.21 (m, 6 H), 7.07 (t, *J* = 7.5 Hz, 1 H), 6.90 – 6.87 (m, 1 H), 6.57 (s, 1 H), 6.22 (d, *J* = 2.4 Hz, 1 H), 5.75 (s, 1 H), 5.45 (dd, *J*₁ = 6.3 Hz, *J*₂ = 2.1 Hz, 1

H), 5.31 (t, $J = 5.1$ Hz, 1 H), 4.92 (dd, $J_1 = 6.6$ Hz, $J_2 = 3.3$ Hz, 1 H), 4.16 – 4.02 (m, 5 H), 3.11 (s, 3 H).

Example 8. 5'-Homologated Derivatives as in Scheme 11:

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3-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-acrylic acid methyl ester:

To a vial containing the starting compound, 1-[9-(6-Hydroxymethyl-2,2-dimethyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl)-9H-purin-6-yl]-3-phenyl-urea (1.07g, 2.5 mmol), in
10 10 ml of dimethylsulfoxide was added IBX (1.06 g, 3.75 mmol) in one portion at RT. The white solid gradually dissolved as the reaction proceeded. After stirring at RT for 2 hours, methyl(triphenylphosphorylidene) acetate (0.84 g, 2.5 mmol) was added in one portion. The reaction was run at RT overnight. To the reaction mixture was added ethyl acetate (100 ml), which was washed with saturated NaHCO_3 , brine, and dried over MgSO_4 . After removal of
15 solvent, the residue was recrystallized with isopropyl alcohol to provide the title compound. MW calculated for $\text{C}_{23}\text{H}_{24}\text{N}_6\text{O}_6$ (MH^+) 481.5 found 481.3 by LCMS.

3-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-propionic acid:

20 To a round bottom flask containing product from the previous step (1.2 g, 2.5 mmol) and palladium on carbon (10 % w/w, 10 mg) under nitrogen was added methanol (20 ml). After flushing with hydrogen gas, the reaction mixture was stirred under a hydrogen atmosphere using a hydrogen balloon overnight at RT. Upon filtration and removal of solvent, the crude white solid product was recrystallized from isopropyl alcohol to give the
25 desired methyl ester compound. MW calculated for $\text{C}_{23}\text{H}_{26}\text{N}_6\text{O}_6$ (MH^+) 483.5 found 483.2.

The methyl ester (500 mg, 1.04 mmol) was dissolved in 4 ml of methanol and sodium hydroxide (83 mg, 2.1 mmol) was then added. The reaction was stirred overnight at RT. After removal of methanol, acetic acid (2.1 mmol, 120 mmol) was added and a white solid precipitated. The solvent was removed under vacuum. The residue was recrystallized from
30 water to provide the pure desired product as a white solid (0.4 g, 83 %). MW calculated for $\text{C}_{22}\text{H}_{24}\text{N}_6\text{O}_6$ (M-1) 467.5 found 467.4 by LCMS.

1-(3-{2,2-Dimethyl-6-[6-(3-phenyl-ureido)-purin-9-yl]-tetrahydro-furo[3,4-d][1,3]dioxol-

4-yl}-propion-1-yl)-pyrrolidine-2-carboxylic acid:

To a vial containing the carboxylic acid product from the previous step (0.117 g, 0.25 mmol), thionyl chloride (0.3 g, 2.5 mmol) was added at 0 °C. After addition, the cold bath was removed followed by addition of two drops of N,N-dimethyl formamide. The reaction mixture was heated to 50 °C for 30 minutes. The excess of thionyl chloride was removed under vacuum and the solid residue was washed with ethyl ether to give the acid chloride. The acid chloride (61 mg, 0.125 mmol) was added to a vial containing L-proline methyl ester (23 mg, 0.138 mmol) and triethylamine (28 mg, 0.275 mmol) in 1 mL of dichloromethane at 0 °C. The reaction was gradually warmed to RT overnight. To the reaction mixture was added ethyl acetate (75 ml), which was washed with 1 N hydrochloric acid, saturated sodium bicarbonate, and dried over magnesium sulfate. The residue was purified by elution from a silica column using 2 % methanol in dichloromethane to give purified prolylmethyl ester product (10 mg, 14 %). MW calculated for C₂₈H₃₃N₇O₇ (MH⁺) 580.6 found 580.3 by LCMS. The prolylmethyl ester (6 mg, 0.010 mmol) was dissolved in tetrahydrofuran (0.1 ml) followed by addition of 6 µl of 15 % sodium hydroxide. After stirring at RT for 2 hrs, ethyl acetate (50 ml) was added and enough 1 N hydrochloric acid was added to adjust the pH to 3. The organic layer was washed with brine and dried over magnesium sulfate. After removal of solvent, the title compound was obtained as a white powder. MW calculated for C₂₇H₃₁N₇O₇ (MH⁺) 566.6 found 566.3 by LCMS.

Example 9. Enzymatic synthesis of a mixture of 1-Ethyl-3-[9-(6-hydroxymethyl-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl)-9H-purin-6-yl]-urea isomers from the corresponding isomeric mixture of 5'-AMP acetal/urea derivatives:

The 2', 3'-(cinnamylacetal)-N⁶-(ethylurea) derivative of AMP (0.750 g, 0.14 mmol) as a mixture of acetal diastereoisomers was dissolved in water (25 mL, 1.4 mol) in a round bottom flask and the pH was adjusted to 8.3 with NaOH. The temperature was adjusted to 35 C, and alkaline phosphatase (0.003 g, 0.00004 mol) was added. Within 15 minutes, the mixture became rather heterogenous, and methanol (20 mL, 0.5 mol) was added to resolubilize the nucleoside product. After 4 h the reaction was judged essentially complete by HPLC. The reaction was worked up by adding more MeOH (20 mL), heating to 60 C to denature the enzyme, and filtering through a 0.22 µm filter. The methanol was evaporated *in vacuo*, and a white, fine-particle solid precipitated from the remaining solvent. This mixture was cooled in an ice bath and filtered. Washing the material with water, followed by drying

over P₂O₅ in a dessicator afforded the title product as a mixture of acetal diastereomers. Dry weight 440 mg (0.10 mmol, 71% yield).

Example 10. 3-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-acrylic acid methyl ester:

1-Ethyl-3-[9-(6-hydroxymethyl-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl)-9H-purin-6-yl]-urea (5.0 g, 11 mmol) was suspended in dry acetonitrile (50 mL) and Dess-Martin periodinate (6.7 g, 16 mmol) was added. The suspension was stirred 2 h, after which time proton NMR of an aliquot showed complete conversion to the aldehyde.

(Methoxycarbonylmethylene)triphenylphosphorane (3.9 g, 12 mmol) was added and stirring was continued overnight. The reaction mixture was then diluted with ethyl acetate (300 mL), washed with saturated sodium bicarbonate/thiosulfate solution (100 mL), dried with sodium sulfate and filtered. The filtrate was evaporated and the solid was dissolved in hot isopropyl alcohol (50 mL). It was allowed to cool, then heptane was added and it was stirred

overnight. The resulting precipitate was washed with heptane and dried under vacuum, affording the desired product (2.9 g, 71%). ¹H-NMR (300 MHz, d₆ DMSO) δ 1.15 (t, 3H, J = 7 Hz), 3.21 (q, 2H, J = 7 Hz), 3.59 (s, 3H), 4.98 (m, 1H), 5.28 (ψt, 1H, J = 6 Hz), 5.51 (dd, 1H, J = 6 Hz, <2 Hz), 5.70 (d, 1H, J = 16 Hz), 5.90 (d, 1H, J = 6 Hz), 6.30 (dd, 1H, J = 6 Hz, 16 Hz), 6.45 (d, 1H, J <2 Hz), 6.95 (d, 1H, J = 15 Hz), 7.35 (m, 3H), 7.45 (d, 2H, J = 7 Hz), 8.50 (s, 1H), 8.60 (s, 1H), 9.30 (t, 1H, J = 6 Hz), 9.60 (s, 1H).

Example 11. 3-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-propionic acid methyl ester:

3-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-acrylic acid methyl ester (250 mg, 0.5 mmol) was dissolved in dry methanol (3 mL). Copper (II) sulfate (90 mg, 0.5 mmol) was added followed by sodium tetrahydroborate (90 mg, 2.5 mmol) and the reaction was stirred 48 h. The reaction was diluted with water, filtered, and concentrated *in vacuo*. The residue was dissolved in ethyl acetate and precipitated with heptane. The precipitate was dissolved in dichloromethane and was chromatographed on silica gel with dichloromethane-methanol (95:5) as eluent, affording the title compound (125 mg, 50%). ¹H-NMR (300 MHz, d₆ DMSO) δ 1.15 (t, 3H, J = 7 Hz), 1.90 (m, 2H), 2.19 (m, 2H), 3.21 (q, 2H, J = 7 Hz), 3.55 (s, 3H), 4.20 (m, 1H), 4.98 (dd, 1H, J = 4 Hz, 6 Hz), 5.45 (dd, 1H, J = 3 Hz, 7 Hz), 5.85 (d, 1H, J = 6 Hz), 6.25 (d, 1H, J = 3 Hz), 6.27 (dd, 1H, J = 6

Hz, 16 Hz), 6.90 (d, 1H, J = 16 Hz), 7.35 (m, 3H), 7.50 (d, 2H, J = 7 Hz), 8.56 (s, 1H), 8.57 (s, 1H), 9.30 (t, 1H, J = 5 Hz), 9.60 (s, 1H).

Example 12. 3-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-propionic acid:

3-{6-[6-(3-Ethyl-ureido)-purin-9-yl]-2-styryl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-propionic acid methyl ester (5.0 g, 10 mmol) was dissolved in tetrahydrofuran (300 mL). Water (100 mL) was added, followed by lithium hydroxide (1.0 g, 25 mmol). The solution was allowed to stir 16 h at room temperature. It was acidified to pH 5 with acetic acid, concentrated *in vacuo*, then extracted with chloroform (300 mL). The organic extract was evaporated, redissolved in ethyl acetate and precipitated with heptane to afford the final product (3.9 g, 80%). ¹H-NMR (300 MHz, d₆ DMSO) δ 1.14 (t 3H, J = 7 Hz), 1.90 (m, 2H), 2.19 (m, 2H), 3.26 (q, 2H, J = 6 Hz), 4.17 (m, 1H), 4.93 (ψt, 1H, J = 6 Hz), 5.43 (dd, 1H, J = 3 Hz, 7 Hz), 5.84 (d 1H, J = 6 Hz), 6.24 (d, 1H, J = 3 Hz), 6.28 (dd, 1H, J = 7 Hz, 13 Hz), 6.90 (d, 1H, J = 16 Hz), 7.35 (m, 3H), 7.51 (d, 2H, J = 7 Hz), 8.56 (s, 1H), 8.61 (s, 1H), 9.30 (t, 1H, J = 6 Hz).

Example 13. Effects of 6-(phenylureido)purine-2', 3'-(4-ethylcyclohexylspiroketyl)-5'-nicotinyl-ribofuranoside (Compound A) and {6-[6-(Cyclohexylmethyl-amino)-purin-9-yl]-2-isobutyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-methanol (Compound B) on the Abdominal Constriction Assay for Visceral Pain in Mice:

This example demonstrates the effects of the above-mentioned two compounds in a model of acetic acid-induced abdominal constriction in mice, a model of visceral pain. The method employed is a modification of the abdominal constriction test described by Collier, *et al.* (Collier, *et al.*, *Br. J. Pharmacol. Chemother.* 32: 295-310 (1968)). For the rest of the example, compound: 6-(phenylureido)purine-2', 3'-(4-ethylcyclohexylspiroketyl)-5'-nicotinyl-ribofuranoside is named "Compound A" and {6-[6-(Cyclohexylmethyl-amino)-purin-9-yl]-2-isobutyl-tetrahydro-furo[3,4-d][1,3]dioxol-4-yl}-methanol is named "Compound B".

Control group pain response

Briefly, a group of eight mice is designated as the control group, and each of said mice in the group receives an intraperitoneal (i.p.) injection of physiological saline in a dose-

volume: mouse-weight ratio of 5 mL/kg ten minutes prior to an i.p. injection of 0.9% acetic acid in physiological saline in a dose-volume: mouse-weight ratio of 10 mL/kg. Each of said mice in the control group is placed in its own observation chamber shortly after the acetic acid injection and is allowed to explore. Each testing period starts 5 minutes after the acetic acid injection into a mouse, and consists of 10 minutes of observation during which time the number of abdominal constrictions of the mouse are counted. Abdominal constriction is defined as a lengthwise stretching of the torso with concave arching of the back. An average number of total constrictions per mouse per observation time is calculated from mice in the control group, and this indicates a baseline level of pain response in the control group animals.

Inhibition of pain response by Compound A

Compound A is administered i.p., at a dose of 50 mg/kg suspended in a volume of physiological saline (5 mL/kg), 10 minutes prior to the administration of acetic acid, as described for the control group above. Eight animals are used in this group and observation of said group demonstrates a statistical difference between the number of averaged abdominal constrictions in the saline control group versus the Compound A group, where the number of averaged abdominal constrictions during the observation time interval is lower when Compound A is introduced, compared to saline control.

Inhibition of pain response by Compound B

Compound B is administered i.p., at a dose of 50 mg/kg suspended in a volume of physiological saline (5 mL/kg), 10 minutes prior to the administration of acetic acid, as described for the control group above. Eight animals are used in this group and observation of said group demonstrates a statistical difference between the number of averaged abdominal constrictions in the saline control group versus the Compound B group, where the number of averaged abdominal constrictions during the observation time interval is lower when Compound B is introduced, compared to saline control.

In similar studies, Compound A and Compound B demonstrate a dose dependent pharmacological effect over saline vehicle control in reducing the abdominal constrictions induced by 0.9% acetic acid. This example demonstrates that compounds of Formula I are effective in treating, preventing or reducing the incidence of visceral pain in animals.

The invention, and the manner and process of making and using it, are now described in such full, clear, concise and exact terms as to enable any person skilled in the art to which it pertains, to make and use the same. It is to be understood that the foregoing describes preferred embodiments of the present invention and that modifications may be made therein without departing from the scope of the present invention as set forth in the claims. To particularly point out and distinctly claim the subject matter regarded as invention, the following claims conclude this specification.

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